## SEARCH REQUEST FORM

## Scientific and Technical Information Center

Art Unit: 1752 Phone N Mail Box and Bldg/Room Location  If more than one search is submi	umber 30 <u>2-13</u> 9069 CRem.) tted, please prid	33 Serial Nu Results Format Pre- pritize searches in	imber:	Date: 1-13-2006 10/815,398 RAPER DISK E-MAIL ed.
********************************  Please provide a detailed statement of the s Include the elected species or structures, ke utility of the invention. Define any terms t known. Please attach a copy of the cover sl	earch topic, and desc ywords, synonyms, hat may have a speci	cribe as specifically as pacronyms, and registry ial meaning. Give exam	oossible the subjournmbers, and comples or relevant	ect matter to be searched.  ombine with the concept or  citations, authors, etc, if
Title of Invention:	Plz. ser	B:6.	SCIE	NTIFIC REFERENCE BE. Sci & rech Inf. Cnt.
Inventors (please provide full names):		-		JAN 1 3 HEL 3
				Pat. & T.M. Office
Earliest Priority Filing Date:			•	Pat. of 1
*For Sequence Searches Only* Please includ appropriate serial number.	e all pertinent informa	tiion (parent, child, divisi	onal, or issued pa	tent numbers) along with the
Plz. Seurch	Bc a	Polymer		
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STAFF USE ONLY	Type of Search		ors and cost who	
Searcher: 111 Le	NA Sequence (#)			
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Searcher Location:	Structure (#)			
Date Searcher Picked Up: 1)8/06  Date Completed: 1)8/06	Bibliographic			·
	Litigation	<del></del>		,
Searcher Prep & Review Time: 60	Fulltext			
Clerical Prep Time: 30	Patent Family			
Online Time: 120	Other	Other (specify)		
PTO-1590 (8-01)	•			

## AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in the application.

Claims 1-3 (Cancelled)

4. (Previously Presented) A compound comprising:

A Polymer comprising:

a polymeric chain;

- an acid labile group attached to the polymeric chain; and

acid de composable at least one hydrophilic group attached to the acid labile group,

acid dissociable

<u>or</u>

wherein the at least one hydrophilic group comprises a sulfhydryl group

Claims 5-14 (Cancelled)

15. (Previously Presented) A compound comprising:

a polymeric chain;

a dissolution inhibitor attached to the polymeric chain to inhibit dissolution of the polymeric chain in a developer; and

solubility sclubility

01

nhibition;

Supplessing

wherein the at least one hydrophilic group comprises a sulfhydryl group

Claims 16-27 (Cancelled)

— SH

- SH



# United States Patent and Trademark Office

UNITED STATES DEPARTMENT OF COMMERCE United States Patent and Trademark Office Address COMMISSIONER FOR PATENTS P.O. Box 1450. Alexandria, Viginia 22313-1450 www.ospto.gov

# \*BIBDATASHEET\*

Bib Data Sheet

**CONFIRMATION NO. 7576** 

SERIAL NUMB 10/815,398		FILING DATE 03/31/2004 RULE	C	CLASS 430	GRO	UP AR1 1752	UNIT		ATTORNEY OCKET NO. 42P18694
APPLICANTS					_				
Wang Yuel	h, Portla	and, OR;							
Ernisse S.	Putna,	Beaverton, OR;							
** CONTINUING : ** FOREIGN APF			* JL *** SJL						
IF REQUIRED, F ** 06/10/2004	ORÈIGI	N FILING LICENSE	GRANTE	ED .					
Foreign Priority claimed 35 USC 119 (a-d) cond met Verified and Acknowledged	ditions (	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	er SJL tials	STATE OR COUNTRY OR	DRA	ETS WING	TOTA	MS	INDEPENDENT CLAIMS 4
ADDRESS 08791 BLAKELY SOKO 12400 WILSHIRE SEVENTH FLOO LOS ANGELES 90025-1030	E BOULI OR	TAYLOR & ZAFMAN EVARD	·						
TITLE Resist compound	is incluc	ding acid labile group	s having	hydrophilic gr	oups at	tached	thereto		
FILING FEE RECEIVED	FEES: / No No	Authority has been gi to charge/cre	iven in Pa edit DEP	aper POSIT ACCOU	NT	1.1 time )	6 Fees (	Proce	essing Ext. of

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=> fil reg
FILE 'REGISTRY' ENTERED AT 13:51:00 ON 18 JAN 2006
=> d his
     FILE 'HCAPLUS' ENTERED AT 11:14:58 ON 18 JAN 2006
L1
              1 S US20050221217/PN
               SEL RN
     FILE 'REGISTRY' ENTERED AT 11:15:33 ON 18 JAN 2006
L2
             5 S E1-E5
     FILE 'LREGISTRY' ENTERED AT 11:55:41 ON 18 JAN 2006
L3
               STR
     FILE 'REGISTRY' ENTERED AT 11:56:51 ON 18 JAN 2006
L4
               SCR 2043
L5
             50 S L3 AND L4
L6
               STR L3
L7
            50 S L6 AND L4
L8
         18969 S L6 AND L4 FUL
                SAV L8 LEE398/A
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L9
         10262 S L8
L10
          47983 S ACID(2A) (DECOMPOS? OR DISSOCIA? OR CLEAVA? OR LABIL?)
L11
            12 S L9(L)L10
L12
            48 S L9 AND L10
L13
            48 S L11 OR L12
L14
        19355 S ?SOLU? (2A) (SUPPRESS? OR INHIBIT?)
L15
             9 S L9 AND L14
L16
            57 S L13 OR L15
L17
           15 S L16 AND PHOTOG?/SC,SX
L18
            8 S L16 AND ?RESIST?
L19
            17 S L17 OR L18
            20 S L16 AND COMPOSITION?
L20
L21
           25 S L19 OR L20
=> d que 121
               SCR 2043
L4
               STR
L6
H \sim S \sim A
3 1 2
NODE ATTRIBUTES:
NSPEC IS RC AT
                      2
CONNECT IS E1 RC AT
DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED
GRAPH ATTRIBUTES:
RING(S) ARE ISOLATED OR EMBEDDED
NUMBER OF NODES IS 3
STEREO ATTRIBUTES: NONE
L8
         18969 SEA FILE=REGISTRY SSS FUL L6 AND L4
L9
         10262 SEA FILE=HCAPLUS ABB=ON PLU=ON L8
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L10
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                DISSOCIA? OR CLEAVA? OR LABIL?)
             12 SEA FILE=HCAPLUS ABB=ON PLU=ON L9(L)L10
L11
L12
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L13
             48 SEA FILE=HCAPLUS ABB=ON PLU=ON L11 OR L12
L14
          19355 SEA FILE=HCAPLUS ABB=ON PLU=ON
                                                  ?SOLU? (2A) (SUPPRESS?
                OR INHIBIT?)
L15
              9 SEA FILE=HCAPLUS ABB=ON PLU=ON L9 AND L14
L16
             57 SEA FILE=HCAPLUS ABB=ON PLU=ON L13 OR L15
             15 SEA FILE=HCAPLUS ABB=ON PLU=ON L16 AND PHOTOG?/SC,SX
L17
L18
              8 SEA FILE=HCAPLUS ABB=ON PLU=ON L16 AND ?RESIST?
             17 SEA FILE=HCAPLUS ABB=ON PLU=ON L17 OR L18
L19
L20
             20 SEA FILE=HCAPLUS ABB=ON
                                         PLU=ON L16 AND COMPOSITION?
             25 SEA FILE=HCAPLUS ABB=ON PLU=ON L19 OR L20
L21
=> fil hcap
FILE 'HCAPLUS' ENTERED AT 13:51:37 ON 18 JAN 2006
=> d l21 1-25 ibib abs hitstr hitind
L21 ANSWER 1 OF 25 HCAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                         2005:1004788 HCAPLUS
DOCUMENT NUMBER:
                         143:315452
TITLE:
                         Resist polymer, resist
                         composition, process for pattern
                          formation, and starting compounds for
                         production of the resist polymer
INVENTOR(S):
                         Momose, Hikaru; Ootake, Atsushi; Nakamura,
                          Tadashi; Ueda, Akifumi
PATENT ASSIGNEE(S):
                         Mitsubishi Rayon Co., Ltd., Japan
SOURCE:
                         PCT Int. Appl., 178 pp.
                          CODEN: PIXXD2
DOCUMENT TYPE:
                         Patent
LANGUAGE:
                          Japanese
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
     PATENT NO.
                         KIND
                                 DATE
                                             APPLICATION NO.
                                                                     DATE
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                          _ _ _ _
     _ _ _ _ _ _ _ _
     WO 2005085301
                          A1
                                 200/50915
                                             WO 2005-JP4402
                                                                     2005
                                                                     0308
             AE, AG, AL, AM, AT/AU, AZ, BA, BB, BG, BR, BW, BY, BZ,
             CA, CH, CN, CO, CP, CU, CZ, DE, DK, DM, DZ, EC, EE, EG,
             ES, FI, GB, GD, 9/E, GH, GM, HR, HU, ID, IL, IN, IS, JP,
             KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL,
             PT, RO, RU, SO, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN,
             TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
         RW: BW, GH, GM, XE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM,
             ZW, AM, AZ,/BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH,
             CY, CZ, DE/ DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT,
             LT, LU, M$\overline{\psi}$, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF,
             CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
PRIORITY APPLN. INFO.:
                                            JP 2004-63616
                                                                     2004
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			0308
JР	2004-73183	Α	
			2004
			0315
JP	2004-189889	Α	
			2004
			0628
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JP	2004-220036	A	
			2004
			0728
TD	2004 252002		
JP	2004-253002	Α	2224
			2004
			0831
qT,	2004-376738	Α	
01	2004 370730	-	2004
			1227
			1227
JP	2005-4315	Α	
			2005
			0111

GI

AB A resist polymer which exhibits reduced line edge roughness and little causes defects in DUV excimer laser lithog. or the like and which contains as the structural unit an acid-dissociable unit represented by the general formula  $J({(K1)k1}_{n}{(L1)11}_{n}{(M1)m1}_{n}{Y}_{n}{(M2)m2}_{n}{(L2)12}_{n}{(K2)k2}$ n[E]n]n or I (n = 2-24; J = n-valent hydrocarbon group optionally bearing a substituent and/or a heteroatom; E = residue of polymerization inhibitor, chain transfer agent, or polymerization initiator; K1 and K2 = at least one kind of groups selected from among alkylene, cycloalkylene, oxyalkylene, arylene, etc.; L1 and L2 = at least one kind of groups selected from among -C(0)0-, -C(0)-, and -OC(0)-; M1, M2, and M3 = at least one kind of groups selected from among alkylene, cycloalkylene, oxyalkylene, and arylene; Y, Y1, and Y2 = acid-dissociable linkage; k1, k2, 11, 12, m1, m2, and m3 = 0 or 1; and R1 is H or Me). 864679-77-6P 864679-79-8P 864683-70-5P IT 864683-73-8P

(preparation of telomer for synthesis of DUV **photoresist** polymer)

RN 864679-77-6 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 3-hydroxytricyclo[3.3.1.13,7]dec-1-yl ester, telomer with 1,1-dimethyl-1,3-propanediyl bis(mercaptoacetate), 2-methyltricyclo[3.3.1.13,7]dec-2-yl 2-methyl-2-propenoate and tetrahydro-2-oxo-3-furanyl 2-methyl-2-propenoate (9CI) (CA INDEX NAME)

CM 1

CRN 864679-76-5 CMF C9 H16 O4 S2

$$\begin{array}{c} \text{O} \\ || \\ \text{O-C-CH}_2\text{--SH} & \text{O} \\ | & || \\ \text{Me-C-CH}_2\text{--CH}_2\text{--O-C-CH}_2\text{--SH} \\ | & \\ \text{Me} \end{array}$$

CM 2

CRN 258879-87-7 CMF (C15 H22 O2 . C14 H20 O3 . C8 H10 O4)x CCI PMS

CM 3

CRN 195000-66-9 CMF C8 H10 O4

CM 4

CRN 177080-67-0 CMF C15 H22 O2

CM 5

CRN 115372-36-6 CMF C14 H20 O3

RN 864679-79-8 HCAPLUS
CN 2-Propenoic acid, 2-methyl-, 3-hydroxytricyclo[3.3.1.13,7]dec-1-yl ester, telomer with 3-mercaptopropanoic acid, 2-methyltricyclo[3.3.1.13,7]dec-2-yl 2-methyl-2-propenoate and tetrahydro-2-oxo-3-furanyl 2-methyl-2-propenoate (9CI) (CA INDEX NAME)

CM 1

CRN 107-96-0 CMF C3 H6 O2 S

 $HS-CH_2-CH_2-CO_2H$ 

CM 2

CRN 258879-87-7 CMF (C15 H22 O2 . C14 H20 O3 . C8 H10 O4)x CCI PMS

CM 3

CRN 195000-66-9 CMF C8 H10 O4

CM 4

CRN 177080-67-0 CMF C15 H22 O2

CRN 115372-36-6 CMF C14 H20 O3

RN 864683-70-5 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 5(or 6)-cyanobicyclo[2.2.1]hept-2-yl ester, telomer with 1-ethylcyclohexyl 2-methyl-2-propenoate and 1,1,4,4-tetramethyl-1,4-butanediyl bis(3-mercaptopropanoate) (9CI) (CA INDEX NAME)

CM 1

CRN 864679-84-5 CMF C14 H26 O4 S2

CM 2

CRN 864683-69-2

CMF (C12 H20 O2 . C12 H15 N O2)x

CCI PMS

CM 3

CRN 274248-09-8 CMF C12 H20 O2

CRN 130668-19-8 CMF C12 H15 N O2 CCI IDS

D1-CN

RN 864683-73-8 HCAPLUS

CN 2-Propenoic acid, 3-hydroxytricyclo[3.3.1.13,7]dec-1-yl ester, telomer with [2-ethyl-2-[[1-[(mercaptoacetyl)oxy]ethoxy]methyl]-1,3-propanediyl]bis(oxyethylidene) bis(mercaptoacetate) and octahydro-1(or 3)-oxo-4,7-methanoisobenzofuran-5-yl 2-propenoate (9CI) (CA INDEX NAME)

CM 1

CRN 864683-72-7 CMF C18 H32 O9 S3

$$\begin{array}{c} \text{O} & \text{O} & \text{O} \\ || & \text{O} \\ \text{CH}_2 - \text{C} - \text{O} & \text{CH}_2 - \text{O} - \text{CH} - \text{Me} \\ || & \text{Me} - \text{CH} - \text{O} - \text{CH}_2 - \text{C} - \text{Et} \\ || & \text{Me} - \text{CH} - \text{O} - \text{CH}_2 \\ || & \text{HS} - \text{CH}_2 - \text{C} - \text{O} \\ || & \text{O} \end{array}$$

CM 2

CRN 864683-71-6

CMF (C13 H18 O3 . C12 H14 O4)x

CCI PMS

CM 3

CRN 436852-35-6 CMF C12 H14 O4

CCI IDS

D2 = 0

CM 4

CRN 216581-76-9 CMF C13 H18 O3

IC ICM C08F020-10

ICS G03F007-039; H01L021-027

CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)
Section cross-reference(s): 35, 38

ST resist DUV photoresist polymer compn telomer chain transfer agent

IT Photolithography

Photoresists

(UV; preparation of telomer for synthesis of DUV photoresist polymer)

IT Chain transfer agents

Resists

(preparation of telomer for synthesis of DUV photoresist polymer)

IT 864679-77-6P 864679-79-8P 864679-81-2P 864683-70-5P 864683-73-8P 864683-74-9P

864683-76-1P 864683-78-3P

(preparation of telomer for synthesis of DUV photoresist polymer)

REFERENCE COUNT:

16 THERE ARE 16 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L21 ANSWER 2 OF 25 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER:

2005:402840 HCAPLUS

DOCUMENT NUMBER:

142:438708

TITLE:

Copolymers for forming fine patterns on

substrates in good adhesiveness, preparation thereof, and thiol compounds therefor

thereof, and thiol compounds therefor Mita, Takahito; Yamagishi, Takanori

INVENTOR(S):
PATENT ASSIGNEE(S):

Maruzen Oil Co., Ltd., Japan

SOURCE:

Jpn. Kokai Tokkyo Koho, 21 pp.

CODEN: JKXXAF

DOCUMENT TYPE:

Patent

LANGUAGE:

Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2005120214	A2	2005Ø5 <u>12</u>	JP 2003-356410	
				2003
PRIORITY APPLN. INFO.:			JP 2003-356410	1016
			01 2003 330110	2003
				1016

OTHER SOURCE(S):

MARAT 142:438708

 $R^{2}$   $R^{3}$   $R^{4}$   $R^{5}$ 

I

AB Ethylenic double bond-containing compds. are radically polymerized using thiol compds. I [R1-R6 = H, Me, Et; ≥1 of R1-R6 = thiol; Y = CH2, (CH2)2, O; n = 0-2] as chain transfer agents to give copolymers (e.g., telomers having terminal groups derived from the thiol compds.), useful for photoresists or antireflective films for semiconductor lithog. The copolymers may comprise (A) units having groups which can be decomposed by acids to increase solubility to alkaline developers and (B) units having polar groups for increase adhesiveness to semiconductor substrates.

IT 851034-01-0P

(thiol compds. as chain transfer agents for preparation of copolymers useful for **photoresists** or antireflective films with good adhesiveness to semiconductor substrates)

RN 851034-01-0 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, hexahydro-2-oxo-3,5-methano-2H-cyclopenta[b]furan-6-yl ester, telomer with dihydro-3-mercapto-2(3H)-furanone and 2-methyltricyclo[3.3.1.13,7]dec-2-yl 2-methyl-2-propenoate (9CI) (CA INDEX NAME)

CM 1

CRN 14032-62-3 CMF C4 H6 O2 S

CM 2

CRN 340964-24-1 CMF (C15 H22 O2 . C12 H14 O4)x CCI PMS

CM 3

CRN 254900-07-7 CMF C12 H14 O4

CM 4

CRN 177080-67-0 CMF C15 H22 O2

IC ICM C08F002-38

ICS C07D307-93; C08F012-00

CC 74-5 (Radiation Chemistry, Photochemistry, and

Photographic and Other Reprographic Processes) Section cross-reference(s): 35, 73, 76 ST butyrolactonyl methacrylate mercaptomethyloxatricyclodecanone telomer photoresist adhesiveness; thiol chain transfer agent telomerization photoresist; semiconductor photolithog antireflective film pos photoresist IT Antireflective films Photolithography Positive photoresists Semiconductor device fabrication Telomerization (thiol compds. as chain transfer agents for preparation of copolymers useful for photoresists or antireflective films with good adhesiveness to semiconductor substrates) IT Telomers (polymers) (thiol compds. as chain transfer agents for preparation of copolymers useful for photoresists or antireflective films with good adhesiveness to semiconductor substrates) IT Chain transfer agents (thiols; thiol compds. as chain transfer agents for preparation of copolymers useful for photoresists or antireflective films with good adhesiveness to semiconductor substrates) IT 851048-16-3P (chain transfer agents; thiol compds. as chain transfer agents for preparation of copolymers useful for photoresists or antireflective films with good adhesiveness to semiconductor substrates) TT 851034-00-9P (in preparation of chain transfer agents; thiol compds. as chain transfer agents for preparation of copolymers useful for photoresists or antireflective films with good adhesiveness to semiconductor substrates) IT 77-73-6, Dicyclopentadiene 591-12-8,  $\alpha$ -Angelical actone (in preparation of chain transfer agents; thiol compds. as chain transfer agents for preparation of copolymers useful for photoresists or antireflective films with good adhesiveness to semiconductor substrates) IT 507-09-5, Thioacetic acid, reactions (in preparation of chain transfer agents; thiol compds. as chain transfer agents for preparation of copolymers useful for photoresists or antireflective films with good adhesiveness to semiconductor substrates) TΥ 851034-01-0P 851048-17-4P (thiol compds. as chain transfer agents for preparation of copolymers useful for photoresists or antireflective films with good adhesiveness to semiconductor substrates) L21 ANSWER 3 OF 25 HCAPLUS COPYRIGHT 2006 ACS on STN ACCESSION NUMBER: 2004:305576 HCAPLUS DOCUMENT NUMBER: 140:347540 TITLE: Negative-working presensitized lithographic plates capable of digital direct laser platemaking INVENTOR(S): Takakuwa, Hideki; Kunita, Kazuto PATENT ASSIGNEE(S): Fuji Photo Film Co., Ltd., Japan SOURCE: Jpn. Kokai Tokkyo Koho, 91 pp. CODEN: JKXXAF

Patent

Japanese

DOCUMENT TYPE:

FAMILY ACC. NUM. COUNT:

LANGUAGE:

#### PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2004117893	A2	20040415	JP/2002-281721	
				2002
				0926
PRIORITY APPLN. INFO.:			/ JP 2002-281721	
				2002
				0926

The lithog. plates comprise, on supports, recording layer containing radical polymerizable monomers LaM(AH)b [L = radical polymerizable group, M = connecting group having valences of (a + b), (AH) = acidic group bearing protonic hydrogen and having acid dissociation constant (pKa) 5.5-13, a = 2-10, b = 1-10], and polymerization initiators. The plates provide scuming-free antiwear printing face.

IT 679843-13-1P 679843-18-4P 679843-20-0P

(polymerized recording layer; neg.-working presensitized lithog. plates having recording layer containing radical polymerizable monomer)

RN 679843-13-1 HCAPLUS

CN Benzoic acid, 2-mercapto-, [4-[[3-[(1-oxooctadecyl)oxy]-2,2-bis[(1-oxo-2-propenyl)oxy]propoxy]carbonyl]phenyl]methyl ester, homopolymer (9CI) (CA INDEX NAME)

CM 1

CRN 679843-04-0 CMF C42 H56 O10 S

RN 679843-16-4 HCAPLUS

CN 2-Propenoic acid, 2-[(3-mercapto-1-oxopropoxy)methyl]-2-[[(1-oxo-2-propenyl)oxy]methyl]-1,3-propanediyl ester, homopolymer (9CI) (CA INDEX NAME)

CM 1

CRN 679843-07-3 CMF C17 H22 O8 S

RN 679843-20-0 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 2,2-bis[(3-mercapto-1-oxopropoxy)methyl]-1,3-propanediyl ester, homopolymer (9CI) (CFINDEX NAME)

CM 1

CRN 679843-10-8 CMF C19 H28 O8 S2

IC ICM G03F007-027 ICS G03F007-00

CC 74-6 (Radiation Chemistry, Photochemistry, and **Photographic** and Other Reprographic Processes) Section cross-reference(s): 38

IT 679843-13-1P 679843-14-2P 679843-15-3P 679843-16-4P 679843-17-5P 679843-18-6P 679843-19-7P 679843-20-0P 679843-21-1P 679843-23-3P

(polymerized recording layer; neg.-working presensitized lithog. plates having recording layer containing radical polymerizable monomer)

L21 ANSWER 4 OF 25 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2003:693498 HCAPLUS

DOCUMENT NUMBER: 139:237456

TITLE: Radiation-sensitive refractive index-variable

compositions for GRIN lens, storage media, and optical device, formation of refractive indexes patterns, and their

patterns

INVENTOR(S): Hanamura, Masaaki; Konno, Keiji; Kumano,

Atsushi; Bessho, Nobuo

PATENT ASSIGNEE(S): JSR Ltd., Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 43 pp.

CODEN: JKXXAF

DOCUMENT TYPE:

Patent Japanese

LANGUAGE:

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2003246930	A2	20030905	JP 2002-48210	
				2002 0225
JP 3632974 PRIORITY APPLN. INFO.:	B2	20050330	JP 2002-48210	
				2002
				0225

AB The compns. contain (A) acid- or basedecomposing polymers having refractive index nA, (B)
nondecomposing compds. having refractive index nB, wherein nA >
nB, selected from ladder polysilsesquioxanes, their hydrolyzates,
and condensates, (C) radiation-sensitive acid or base generator,
and (D) stabilizers. Thus, a composition comprising
poly(o-phthalaldehyde) 40, GR 650 (B) 60, 2-(4-methoxyphenyl)-4,6bis(trichloromethyl)-s-triazine 1 part, thinned in diethylene
glycol Me Et ether, was applied on a Si wafer, pre-baked to give a
film, exposed to light (λ = 365 nm), and subjected to PEB
(post-exposure-bake) to give a 3.0-μm thick refractive indexes
pattern having regions with refractive index 1.42 and 1.52.

IT 402739-88-2P 402739-89-3P 402739-90-6P

, 1,4-Benzenedithiol-p-phenylene diisocyanate copolymer 402739-92-8P, 1,4-Benzenedithiol-4-nitro-1,3-phenylenedichloroformate copolymer

(radiation-sensitive refractive index-variable compns

. and formation of refractive indexes patterns for GRIN lens, storage media, and optical device)

RN 402739-88-2 HCAPLUS

CN 1,4-Benzenedithiol, polymer with 1,4-bis(2-nitroethenyl)benzene (9CI) (CA INDEX NAME)

CM 1

CRN 3179-01-9 CMF C10 H8 N2 O4

$$O_2N-CH$$
 CH  $CH-NO_2$ 

CM 2

CRN 624-39-5 CMF C6 H6 S2

RN 402739-89-3 HCAPLUS

CN 1,4-Benzenedicarbonyl dichloride, polymer with 1,4-benzenedithiol (9CI) (CA INDEX NAME)

CM 1

CRN 624-39-5 CMF C6 H6 S2

CM 2

CRN 100-20-9 CMF C8 H4 Cl2 O2

RN 402739-90-6 HCAPLUS

CN 1,4-Benzenedithiol, polymer with 1,4-diisocyanatobenzene (9CI) (CA INDEX NAME)

CM 1

CRN 624-39-5 CMF C6 H6 S2

CM 2

CRN 104-49-4 CMF C8 H4 N2 O2

RN 402739-92-8 HCAPLUS

CN Carbonochloridic acid, 4-nitro-1,3-phenylene ester, polymer with 1,4-benzenedithiol (9CI) (CA INDEX NAME)

CM 1

CRN 402739-91-7 CMF C8 H3 Cl2 N O6

CM 2

CRN 624-39-5 CMF C6 H6 S2

IC ICM C08L101-00

ICS C08K005-00; C08L083-04; G02B001-04; G02B003-00; G02B005-18; G02B006-00; G02B006-12; G03F007-004; G03H001-02

CC 73-11 (Optical, Electron, and Mass Spectroscopy and Other Related Properties)

Section cross-reference(s): 38

radiation sensitive refractive index variable compn;
GRIN lens polysilsesquioxane decompg polymer compn;
hologram device polysilsesquioxane decompg polymer compn;
diffraction grating polysilsesquioxane decompg polymer compn; optical fiber polysilsesquioxane decompg polymer compn; optical disk polysilsesquioxane decompg polymer compn; optical integrated circuit polysilsesquioxane decompg polymer compn

IT Silsesquioxanes

(Me Ph, Glass Resin GR 908, GR 950; radiation-sensitive refractive index-variable compns. and formation of

refractive indexes patterns for GRIN lens, storage media, and optical device) TT Silsesquioxanes (Me Ph, ethoxy- and hydroxy-terminated, Glass Resin GR 100; radiation-sensitive refractive index-variable compns. and formation of refractive indexes patterns for GRIN lens, storage media, and optical device) ITSilsesquioxanes (Me, Glass Resin GR 650; radiation-sensitive refractive index-variable compns. and formation of refractive indexes patterns for GRIN lens, storage media, and optical device) Integrated circuits IT (optical; radiation-sensitive refractive index-variable compns. and formation of refractive indexes patterns for GRIN lens, storage media, and optical device) TΤ Diffraction gratings Holography Optical disks Optical instruments (radiation-sensitive refractive index-variable compns . and formation of refractive indexes patterns for GRIN lens, storage media, and optical device) Polycarbonates, uses TT Polyethers, uses (radiation-sensitive refractive index-variable compns . and formation of refractive indexes patterns for GRIN lens, storage media, and optical device) IT Polyurethanes, uses (thio-; radiation-sensitive refractive index-variable compns. and formation of refractive indexes patterns for GRIN lens, storage media, and optical device) ΙT 3584-23-4, 2-(4-Methoxyphenyl)-4,6-bis(trichloro methyl)-s-triazine 16668-99-8, Diphenyliodonium trifluoroacetate 111281-12-0 (radiation-sensitive acid generator; radiation-sensitive refractive index-variable compns. and formation of refractive indexes patterns for GRIN lens, storage media, and

optical device) TΤ 119313-12-1, 2-Benzyl-2-dimethylamino-1-(4-morpholinophenyl)-butan-204782-35-4, N-(2-Nitrobenzyloxycarbonyl)pyrrolidine (radiation-sensitive base generator; radiation-sensitive refractive index-variable compns. and formation of refractive indexes patterns for GRIN lens, storage media, and

IT 643-79-8DP, o-Phthalaldehyde, reaction products with potassium tert-butoxide 865-47-4DP, reaction products with o-phthalaldehyde

optical device)

IT

(radiation-sensitive refractive index-variable compns . and formation of refractive indexes patterns for GRIN lens, storage media, and optical device)

25750-62-3P, o-Phthalaldehyde homopolymer 26966-00-7P 119652-66-3P 119806-22-3P 155462-47-8P 155462-48-9P 402739-87-1P 402739-88-2P 402739-89-3P 402739-90-6P, 1,4-Benzenedithiol-p-phenylene diisocyanate copolymer 402739-92-8P, 1,4-Benzenedithiol-4-nitro-1,3phenylenedichloroformate copolymer 402739-96-2P 402739-97-3P, 1,4-Benzenedithiol-p-phenylene diisocyanate copolymer, sru 402745-46-4P 402859-28-3P 592537-96-7P 592538-05-1P 592538-08-4P 592538-11-9P

```
(radiation-sensitive refractive index-variable compns
        . and formation of refractive indexes patterns for GRIN lens,
        storage media, and optical device)
     122-60-1, Phenyl glycidyl ether 539-48-0, 1,4-Benzenedimethanamine 1675-54-3, Bisphenol A diglycidyl ether
IT
     3221-14-5, 3-Phenoxypropylene sulfide 7426-75-7
         (stabilizer; radiation-sensitive refractive index-variable
        compns. and formation of refractive indexes patterns
        for GRIN lens, storage media, and optical device)
L21 ANSWER 5 OF 25 HCAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                          2003:376854 HCAPLUS
DOCUMENT NUMBER:
                          138:386271
TITLE:
                          Thermally dissociable maleic
                          acid derivative crosslinker and
                          curable composition containing the
                          same
INVENTOR(S):
                          Choi, Wonmun
PATENT ASSIGNEE(S):
                          The Yokohama Rubber Co., Ltd., Japan,
                          PCT Int. Appl., 50 pp.
SOURCE:
                          CODEN: PIXXD2
DOCUMENT TYPE:
                          Patent
LANGUAGE:
                          Japanese
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
     PATENT NO.
                          KIND
                                               APPLICATION NO.
                                  DATE
                                                                       DATE
     -----
     _____
                                               WO 2002-JP11369
     WO 2003040133
                           A1
                                  20030515
                                                                        2002
                                                                        1031
             AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA,
              CH, CN, CO, CR, CU, CZ, DE, DK, PM, DZ, EC, EE, ES, FI,
             GB, GD, GE, GH, GM, HR, HU, ID, /IL, IN, IS, JP, KE, KG,
             KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE,
             SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ,
             VC, VN, YU, ZA, ZM, ZW
         RW: GH, GM, KE, LS, MW, MZ, SD/SL, SZ, TZ, UG, ZM, ZW, AT,
             BE, BG, CH, CY, CZ, DE, DX, EE, ES, FI, FR, GB, GR, IE,
              IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF, CG, CI, CM,
             GA, GN, GQ, GW, ML, MR,/NE, SN, TD, TG
     US 2004260038
                           A1
                                  200/1223
                                              US 2004-493232
                                                                        2004
                                                                        0421
PRIORITY APPLN. INFO.:
                                               JP 2001-340779
                                                                        2001
                                                                        1106
                                               JP 2001-381198
                                                                        2001
                                                                        1214
                                               JP 2001-381199
                                                                        2001
                                                                       1214
                                               WO 2002-JP11369
                                                                       2002
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OTHER SOURCE(S): MARPAT 138:386271

AB The maleic acid deriv.is obtained by reacting a compound having ≥1 thiol groups with a maleic acid compound The maleic acid derivative is thermally dissociable. The curable composition, which contains this maleic acid derivative as a hardener, is usable as a thermally dissociable material. Reacting 2-di-n-butylamino-4,6-dimercapto-s-triazine with maleic anhydride in MEK at 70° for 3 h, mixing the resulting adduct with bisphenol A type epoxy resin and DMP 30, and curing 30 min at 80° gave a tack-free product, which was thermally dissociated when heated in a 160° oven for 10 min.

IT 525601-96-1P

(oligomeric, crosslinker; thermally dissociable
maleic acid derivative crosslinker for curable
composition with good disposability)

RN 525601-96-1 HCAPLUS

CN 1H-Pyrrole-2,5-dione, 1,1'-(1,2-ethanediyl)bis-, polymer with 6-(dibutylamino)-1,3,5-triazine-2,4(1H,3H)-dithione (9CI) (CA INDEX NAME)

CM 1

CRN 29529-99-5 CMF C11 H20 N4 S2

CM 2

CRN 5132-30-9 CMF C10 H8 N2 O4

IC ICM C07D405-14 ICS C07D417-14; C07D251-46; C07D207-416; C07D403-14; C08G075-00; C08G018-38; C08G059-40

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CC
     37-3 (Plastics Manufacture and Processing)
     Section cross-reference(s): 27
ST
     thermally dissociable maleic acid deriv
     crosslinker; maleic anhydride dibutylaminodimercaptotriazine
     adduct crosslinker epoxy resin
IT
     Polyurethanes, uses
        (polyoxyalkylene-, thermally dissociable; thermally
        dissociable maleic acid derivative crosslinker
        for curable composition with good disposability)
IT
     Crosslinking agents
     Polymer degradation
        (thermally dissociable maleic acid derivative
        crosslinker for curable composition with good
        disposability)
IT
     Epoxy resins, uses
        (thermally dissociable; thermally dissociable maleic
        acid derivative crosslinker for curable composition
        with good disposability)
IT
     525601-94-9P
                    525601-95-0P
                                   525601-97-2P
                                                  525601-98-3P
     526211-13-2P
        (crosslinker; thermally dissociable maleic
        acid derivative crosslinker for curable composition
        with good disposability)
IT
     525601-96-1P
        (oligomeric, crosslinker; thermally dissociable
        maleic acid derivative crosslinker for curable
        composition with good disposability)
IT
     9057-91-4, Polypropylene glycol-TDI copolymer
     Bisphenol A epoxy resin 502476-12-2, ED505R
        (thermally dissociable maleic acid derivative
        crosslinker for curable composition with good
        disposability)
TΤ
     60-24-2, 2-Mercaptoethanol 108-31-6, Maleic anhydride, reactions
     147-93-3, Thiosalicylic acid 925-21-3, Maleic acid
     Monobutyl ester 1072-71-5, 1,3,4-Thiadiazole-2,5-dithiol
     5132-30-9, 1,2-Bis (maleimido) ethane
                                           29529-99/-5,
     2-Dibutylamino-4,6-dimercapto-s-triazine
        (thermally dissociable maleic acid derivative
        crosslinker for curable composition with good
        disposability)
                               THERE ARE 10 CITED REFERENCES AVAILABLE
REFERENCE COUNT:
                         10
                               FOR THIS RECORD. ALL CITATIONS AVAILABLE
                               IN THE RE FORMAT
L21 ANSWER 6 OF 25 HCAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                         2003:94105 ACAPLUS
DOCUMENT NUMBER:
                         138:156904
TITLE:
                         Easily decomposable corrosion inhibitor for
                         metal
                         Oda, Yoshihisa
INVENTOR(S):
                         Nippon/Shokubai Co., Ltd., Japan
PATENT ASSIGNEE(S):
SOURCE:
                         Jpn. Kokai Tokkyo Koho, 15 pp.
                         CODEN: JKXXAF
DOCUMENT TYPE:
                         Patent
LANGUAGE:
                         Japanese
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
     PATENT NO.
                         KIND
                                DATE
                                           APPLICATION NO.
                                                                   DATE
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JP 2003034677

A2 20030207

JP 2001-217584

2001 0718

PRIORITY APPLN. INFO.:

JP 2001-217584

2001 0718

OTHER SOURCE(S): MARPAT 138:156904

AB The inhibitor comprises [HS(CnHm)OCO]zCiHj(CO2M)k [I; n = 1-12; m = 2-2n; i = 2-25; j = 0-(2i+1-k-z); k = 1-5; z = 1-5; M = H, metal, ammonium]. Corrosion inhibitor compns. comprise salts and/or mixts. of I and polyamines or [HS(CnHm)OCO]zCiHj(CO2H)k.1/sR(CtNuNv)wR' [n, m, i, j, k, z = same as I; t = 2-10, u = 5-25; v = 1-5; w = 2-200,000; v = 10, amino, C1-20 hydrocarbyl; v = 1-10 (v = 11).

IT 494825-09-1 494825-10-4

(easily decomposable mercaptooxycarbonylcarboxylic
acid corrosion inhibitor for metal)

RN 494825-09-1 HCAPLUS

CN 1,2-Benzenedicarboxylic acid, mono(2-mercaptoethyl) ester, polymer
with aziridine (9CI) (CA INDEX NAME)

CM 1

CRN 128338-14-7 CMF C10 H10 O4 S

CM 2

CRN 151-56-4 CMF C2 H5 N

H N

RN 494825-10-4 HCAPLUS

CN 1,2,3-Benzenetricarboxylic acid, 1-(2-mercaptoethyl) ester, polymer with aziridine (9CI) (CA INDEX NAME)

CM 1

CRN 142495-88-3 CMF C11 H10 O6 S

```
CO<sub>2</sub>H |
HO<sub>2</sub>C
                - o- сн<sub>2</sub>- сн<sub>2</sub>- sн
     CM
          2
     CRN 151-56-4
     CMF C2 H5 N
ΙÇ
     ICM C07C323-12
     ICS C23F011-00; C23F011-16
CC
     55-10 (Ferrous Metals and Alloys)
     Section cross-reference(s): 23, 56
TT
     Corrosion inhibitors
        (easily decomposable mercaptooxycarbonylcarboxylic
        acid corrosion inhibitor for metal)
IT
     9002-98-6D, reaction products with dicarboxylic acid
     monomercaptoalkyl ester
                               128338-14-7
                                               142495-88-3
     494825-09-1 494825-10-4
                               494836-07-6
        (easily decomposable mercaptooxycarbonylcarboxylic
        acid corrosion inhibitor for metal)
IT
     7429-90-5, Aluminum, uses
                                 7439-89-6, Iron, uses
                                                            39462-15-2,
     uses
        (easily decomposable mercaptooxycarbonylcarboxylic)
        acid corrosion inhibitor for metal)
ΙT
     25377-73-5, Dodecenylsuccinic anhydride
        (in preparation of inhibitor; easily decomposable
        mercaptooxycarbonylcarboxylic acid corrosion
        inhibitor for metal)
L21 ANSWER 7 OF 25 HCAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                          2002:975711 HCAPLUS
DOCUMENT NUMBER:
                          138:63820
TITLE:
                          Polybenzoxazole precursors, their preparation,
                          and positive photoresists therewith
INVENTOR(S):
                          Ehara, Kazuya; Shibazaki, Yuji; Ueda, Mitsuru
PATENT ASSIGNEE(S):
                          Nitto Denko Corp. / Japan
SOURCE:
                          Jpn. Kokai Tokkyø Koho, 9 pp.
                          CODEN: JKXXAF
DOCUMENT TYPE:
                          Patent
LANGUAGE:
                          Japanese
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
     PATENT NO.
                          KIND
                                 DATE
                                              APPLICATION NO.
                                                                      DATE
     JP 2002371133
                           A2
                                 20021226
                                              JP 2001-180679
```

2001

PRIORITY APPLN. INFO.:

JP 2001-180679

2001

0614

0614

OTHER SOURCE(S): MARPAT 138:63820

AB Active carboxylic acids XCOR1COX [R1 = bivalent organic group; X = leaving group (e.g., cyclic compound residues containing N, S, C, O, and aromatic ring)] and dihydroxydiamines R2(NH2)2(OH)2 (R2 = tetravalent organic group) are reacted in organic solvents to afford polybenzoxazole precursors which can be directly (i.e., without isolation) formulated with other photoresist components.

IT 479198-42-0P

(preparation of polybenzoxazole precursors from carboxylic acids and dihydroxydiamines for pos. **photoresists**)

RN 479198-42-0 HCAPLUS

CN Poly[iminocarbonothioyl-1,3-phenylenecarbonothioylimino(6-hydroxy-1,3-phenylene)[2,2,2-trifluoro-1-(trifluoromethyl)ethylidene](4-hydroxy-1,3-phenylene)] (9CI) (CA INDEX NAME)

IC ICM C08G073-22

ICS G03F007-023; G03F007-032

CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)
Section cross-reference(s): 35

ST polybenzoxazole precursor prepn isophthaloylbisbenzoxazinethione aminohydroxyphenylfluoropropane reaction; photoresist polybenzoxazole precursor polyhydroxyamide synthesis

IT Polybenzoxazoles

(fluorine-containing; preparation of polybenzoxazole precursors from carboxylic acids and dihydroxydiamines for pos.

photoresists)
Fluoropolymers, preparation

(polybenzoxazole-; preparation of polybenzoxazole precursors from carboxylic acids and dihydroxydiamines for pos. photoresists).

IT Positive photoresists

(preparation of polybenzoxazole precursors from carboxylic acids and dihydroxydiamines for pos. **photoresists**)

IT 479198-43-1

IT

(dissoln. inhibitors; preparation of polybenzoxazole precursors from carboxylic acids and dihydroxydiamines for pos. photoresists)

IT 80292-82-6P

(in preparation of polybenzoxazole precursors from carboxylic acids and dihydroxydiamines for pos. photoresists)

IT 99-63-8, Isophthaloyl chloride 149-30-4, 2-Mercaptobenzothiazole

(in preparation of polybenzoxazole precursors from carboxylic acids

2382-96-9, 2-Mercaptobenzoxazole

```
and dihydroxydiamines for pos. photoresists)
ΙT
     113339-21-2P
                    370567-91-2P 479198-42-0P
        (preparation of polybenzoxazole precursors from carboxylic acids and
        dihydroxydiamines for pos. photoresists)
L21 ANSWER 8 OF 25 HCAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                         2001:900258 HCAPLUS
DOCUMENT NUMBER:
                         136:29177
TITLE:
                         Radiation-sensitive resin composition
                         for chemical amplified pos. tone resist
INVENTOR(S):
                         Nishimura, Yukio; Douki, Katsuji; Kajita,
                         Toru; Shimokawa, Tsutomu
PATENT ASSIGNEE(S):
                         JSR Corporation, Japan
                         Eur. Pat. Appl., 54 pp.
SOURCE:
                         CODEN: EPXXDW
DOCUMENT TYPE:
                         Patent
LANGUAGE:
                         English
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
                                             APPLICATION NO.
     PATENT NO.
                         KIND
                                 DATE
                                                                     DATE
     EP 1162506
                                 20011212
                                                2001-113944
                          A1
                                                                     2001
                                                                     0607
         R: AT, BE, CH, DE, DK, ES, FR, GP,
                                             GR, IT, LI, LU, NL, SE,
             MC, PT, IE, SI, LT, LV, FI, KO
     JP 2002062657
                          A2
                                 20020228/
                                             JP 2001-95877
                                                                     2001
                                                                     0329
                                 20020124
     US 2002009667
                          A1
                                             US 2001-874977
                                                                     2001
                                                                     0607
     US 6753124
                                 200/40622
                          B2
PRIORITY APPLN. INFO.:
                                             JP 2000-173708
                                                                     2000
                                                                     0609
                                             JP 2001-95877
                                                                     2001
                                                                     0329
GI
(R^1)b
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AB A radiation-sensitive resin composition used as a chemical amplified pos. tone resist responsive to short wavelength active radiation such as KrF excimer laser and ArF excimer laser is disclosed. The resin composition comprises: (A) an acid-dissociable group-containing resin which is insol. or scarcely soluble in alkali and becomes alkali soluble when the acid-dissociable group dissocs., the resin comprising a lactone cyclic structure I (a = 1-3; b = 0-9; R1 = monovalent organic group); and (B) aphotoacid generator. The composition has high transmittance of radiation, exhibits high sensitivity, resolution, and pattern shape, and can produce semiconductors at a high yield without producing resolution defects during microfabrication. IT

379257-71-3P

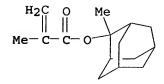
(radiation-sensitive resin composition for chemical amplified pos. tone resist)

RN 379257-71-3 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 2-methyltricyclo[3.3.1.13,7]dec-2-yl ester, polymer with 3-mercaptopropanoic acid and (tetrahydro-5-oxo-2-furanyl)methyl 2-methyl-2-propenoate (9CI) (CA INDEX NAME)

CM 1

CRN 177080-67-0 CMF C15 H22 O2



CM 2

CRN 156938-09-9 CMF C9 H12 O4

$$\begin{array}{c|c} \text{O} & \text{CH}_2 \\ \text{O} & \text{CH}_2 - \text{O} - \text{C} - \text{C} - \text{Me} \end{array}$$

CM 3

CRN 107-96-0 CMF C3 H6 O2 S

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HS-CH_2-CH_2-CO_2H
TC
     ICM G03F007-039
     74-5 (Radiation Chemistry, Photochemistry, and
CC
     Photographic and Other Reprographic Processes)
     Section cross-reference(s): 35, 38, 76
ST
     chem amplified photoresist acid
     dissociable lactone cyclic
     Photolithography
IT
        (UV; radiation-sensitive resin composition for)
IT
     Positive photoresists
        (radiation-sensitive resin composition for)
TT
     1116-76-3, Tri-n-octylamine
                                  3033-62-3, Bis(2-
     dimethylaminoethyl)ether
                               193810-83-2, N-tert-Butoxycarbonyl-2-
     phenylbenzimidazole
                           330576-56-2, N-tert-
     Butoxycarbonyldicyclohexylamine
        (acid diffusion control agent; radiation-sensitive resin
        composition for chemical amplified pos. tone resist)
IT
     157692-53-0, tert-Butyl deoxycholate
                                           169228-97-1, Di-tert-butyl
     1,3-adamantanedicarboxylate 231296-44-9, t-Butoxycarbonylmethyl
                    296242-01-8, 2,5-Dimethyl-2,5-di(1-
     deoxycholate
     adamantylcarbonyloxy) hexane
        (additives; radiation-sensitive resin composition for
        chemical amplified pos. tone resist)
TT
     542-92-7, Cyclopentadiene, reactions
                                           814-68-6, Acryloyl chloride
     920-46-7, Methacryloyl chloride 34862-06-1, 4,5-
     Diacetoxypentanal
                         78984-88-0
        (preparation of radiation-sensitive resin composition for chemical
        amplified pos. tone resist)
IT
     10374-51-3P, 4-Hydroxymethyl-γ-butyrolactone
                                                    156938-09-9P
     259154-20-6P, 4,5-Diacetoxypentanoic acid 264193-11-5P
     379257-66-6P
                    379257-68-8P
                                   379257-69-9P
                                                  379257-70-2P
        (preparation of radiation-sensitive resin composition for chemical
        amplified pos. tone resist)
IT
     379257-71-3P
                    379257-72-4P
                                   379257-73-5P
                                                  379257-75-7P
     379257-76-8P
                    379257-77-9P
                                   379257-78-0P
                                                  379257-79-1P
     379257-81-5P
                    379257-82-6P
                                   379257-83-7P
        (radiation-sensitive resin composition for chemical amplified
       pos. tone resist)
IT
     96-48-0, γ-Butyrolactone
                                108-94-1, Cyclohexanone, uses
     110-43-0, 2-Heptanone 84540-57-8, Propylene glycol monomethyl
     ether acetate
        (solvent; radiation-sensitive resin composition for chemical
        amplified pos. tone resist)
REFERENCE COUNT:
                               THERE ARE 4 CITED REFERENCES AVAILABLE
                         4
                               FOR THIS RECORD. ALL CITATIONS AVAILABLE
                               IN THE RE FORMAT
L21 ANSWER 9 OF 25 HCAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                         2001:736888 HCAPLUS
DOCUMENT NUMBER:
                         135:290228
TITLE:
                         Manufacture of water-based emulsions
                         containing self-emulsifying actylic urethane
                         polymers for coatings
INVENTOR(S):
                         Morishima, Takeshi; Kanno, Akira; Sasahara,
                         Toshiaki; Sugimoto, Kazuya; Shimura, Noriyuki;
                         Yamaquchi, Hirotaka
PATENT ASSIGNEE(S):
                         Nippon Polyurethane Industry Co., Ltd., Japan;
                          USHA SHRESTHA EIC 1700 REM 4B28
```

Asia Industry Co., Ltd.

SOURCE: Jpn. Kokai Tokkyo Koho, 21 pp.

CODEN: JKXXAF

DOCUMENT TYPE: LANGUAGE:

Patent Japanese

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2001278923	A2	20011010	JP 2000-98184	
				2000
				0331
PRIORITY APPLN. INFO.:			JP 2000-98184	
				2000
				0331

AB The emulsions with good storage stability for haze-free coatings with high flexibility, toughness, durability, and solvent resistance, are manufactured without using organic solvents by (1) reacting organic polyisocyanates with (A) SH- and active H-containing compds., (B) active H-containing compds. having hydrophilic anion- or cation-forming groups, and (C) other active H-containing compds. in ethylenically unsatd. monomers in the presence of polymerization inhibitors' to give solns. of active H-terminated urethane prepolymers having SH, hydrophilic anion- or cation-forming groups, and number-average mol. weight ≤50,000 in the unsatd. monomers, (2) neutralizing the solns. with cations or cationic neutralization agents, (3) emulsifying the solns. in H2O, and (4) radically polymerizing the urethane prepolymers with the unsatd. monomers in the presence of radical polymerization initiators. Alternatively, the emulsions are manufactured by preparing solns. of urethane prepolymers similarly without using A, neutralizing the solns., adding H2O, aminothiols, and optionally chain extenders to the neutralized solns. to form emulsions of the SH-containing prepolymers, and carrying out the above radical polymerization prepolymer prepared by reacting polyester diol (prepared from 3-methyl-1,5-pentanediol and adipic acid), neopentyl glycol, 2,2-dimethylol butyric acid, 2-mercapto-1-hydroxyethane, and 1,6-hexamethylene diisocyanate in Me methacrylate, was mixed with acrylic acid, diacetoneacrylamide, Me methacrylate, and Bu acrylate, neutralized with triethylamine, emulsified with H2O, radical-polymerized, and reacted with qlutaric acid dihydrazide to give an aqueous emulsion, which was used to form a film showing high water resistance.

IT 365438-66-0P 365438-68-2P 365438-70-6P 365438-72-8P 365438-74-0P 365438-77-3P 365438-79-5P 365438-81-9P 365438-83-1P 365438-85-3P 365438-88-6P

(manufacture of aqueous acrylic urethane polymer emulsions without using organic solvents for haze-free flexible coatings)

RN 365438-66-0 HCAPLUS

CN Hexanedioic acid, polymer with 2,2-bis(hydroxymethyl)butanoic acid, butyl 2-propenoate, 1,6-diisocyanatohexane, N-(1,1-dimethyl-3-oxobutyl)-2-propenamide, 2,2-dimethyl-1,3-propanediol, 2-mercaptoethanol, methyl 2-methyl-2-propenoate, 3-methyl-1,5-pentanediol, pentanedioic acid dihydrazide and 2-propenoic acid, compd. with N,N-diethylethanamine (9CI) (CA INDEX NAME)

CRN 121-44-8 CMF C6 H15 N

Et | Et-N-Et

CM 2

CRN 365438-65-9

CMF (C9 H15 N O2 . C8 H12 N2 O2 . C7 H12 O2 . C6 H14 O2 . C6 H12 O4 . C6 H10 O4 . C5 H12 N4 O2 . C5 H12 O2 . C5 H8 O2 . C3 H4 O2 . C2 H6 O S)  $\times$ 

CCI PMS

CM 3

CRN 10097-02-6 CMF C6 H12 O4

 $_{\rm CH_2-OH}^{\rm CH_2-OH}$   $_{\rm CH_2-OH}^{\rm CH_2-OH}$ 

CM 4

CRN 4457-71-0 CMF C6 H14 O2

 $\begin{array}{c} & \text{Me} \\ | \\ \text{HO-} \ \text{CH}_2 - \text{CH}_2 - \text{CH-} \ \text{CH}_2 - \text{CH}_2 - \text{OH} \end{array}$ 

CM 5

CRN 2873-97-4 CMF C9 H15 N O2

$$\begin{array}{c} \text{O} & || \\ \text{H}_2\text{C} = \text{CH} - \text{C} - \text{NH} & \text{O} \\ | & || \\ \text{Me} - \text{C} - \text{CH}_2 - \text{C} - \text{Me} \\ | & \\ \text{Me} \end{array}$$

CRN 1508-67-4 CMF C5 H12 N4 O2

$$\begin{array}{c|c} & \circ & \circ \\ \parallel & \parallel \\ \text{$_{12}$N-NH-C- (CH$_{2})$} & \text{$_{3}$-C-NH-NH$_{2}$} \end{array}$$

CM 7

CRN 822-06-0 CMF C8 H12 N2 O2

$$OCN-(CH2)6-NCO$$

CM 8

CRN 141-32-2 CMF C7 H12 O2

CM 9

CRN 126-30-7 CMF C5 H12 O2

CM 10

CRN 124-04-9 CMF C6 H10 O4

 $HO_2C^-$  (CH<sub>2</sub>)<sub>4</sub>-CO<sub>2</sub>H

CM 11

CRN 80-62-6 CMF C5 H8 O2

CM 12

CRN 79-10-7 CMF C3 H4 O2

0 || но- с- сн== сн<sub>2</sub>

CM 13

CRN 60-24-2 CMF C2 H6 O S

 $HO-CH_2-CH_2-SH$ 

RN 365438-68-2 HCAPLUS
CN Butanoic acid, 2,2-bis(hydroxymethyl)-, polymer with
bis(isocyanatomethyl)cyclohexane, butyl 2-propenoate, diethyl
carbonate, 1,3-dihydroxy-2-propanone, 2,3-dimercapto-1-propanol,
N-(1,1-dimethyl-3-oxobutyl)-2-propenamide, α,α'-1,2ethanediylbis[ω-hydroxypoly[oxy(1-oxo-1,6-hexanediyl)]],
ethenylbenzene, 1,6-hexanediol, 2-hydroxyethyl
2-methyl-2-propenoate, 2-mercaptoethanol, methyl
2-methyl-2-propenoate, pentanedioic acid dihydrazide and
2-propenoic acid, compd. with N,N-diethylethanamine (9CI) (CA
INDEX NAME)

CM 1

CRN 121-44-8 CMF C6 H15 N

CRN 365438-67-1

CMF (C10 H14 N2 O2 . C9 H15 N O2 . C8 H8 . C7 H12 O2 . C6 H14 O2 . C6 H12 O4 . C6 H10 O3 . (C6 H10 O2)n (C6 H10 O2)n C2 H6 O2 . C5 H12 N4 O2 . C5 H10 O3 . C5 H8 O2 . C3 H8 O S2 . C3 H6 O3 . C3 H4 O2 . C2 H6 O S)x

CCI PMS

CM 3

CRN 59692-54-5 CMF (C6 H10 O2)n (C6 H10 O2)n C2 H6 O2 CCI PMS

CM 4

CRN 42170-25-2 CMF C10 H14 N2 O2 CCI IDS



$$2 \left\lceil D1 - CH_2 - NCO \right\rceil$$

CM 5

CRN 10097-02-6 CMF C6 H12 O4

CRN 2873-97-4 CMF C9 H15 N O2

$$\begin{array}{c} {\rm O} \\ || \\ {\rm H_2C} = {\rm CH-C-NH} \\ || \\ {\rm Me-C-CH_2-C-Me} \\ || \\ {\rm Me} \end{array}$$

CM 7

CRN 1508-67-4 CMF C5 H12 N4 O2

$$\begin{array}{c|c} & & & \circ & \\ \parallel & & \parallel & \\ \text{H}_2\text{N}-\text{NH}-\text{C}-\text{(CH}_2)}_3-\text{C}-\text{NH}-\text{NH}_2 \\ \end{array}$$

CM 8

CRN 868-77-9 CMF C6 H10 O3

CM 9

CRN 629-11-8 CMF C6 H14 O2

 $_{\rm HO^-}$  (CH<sub>2</sub>)<sub>6</sub>-OH

CM 10

CRN 141-32-2 CMF C7 H12 O2

CRN 105-58-8 CMF C5 H10 O3

CM 12

CRN 100-42-5 CMF C8 H8

$$H_2C = CH - Ph$$

CM 13

CRN 96-26-4 CMF C3 H6 O3

$$_{\rm HO-\,CH_2-C-\,CH_2-\,OH}^{\rm O}$$

CM 14

CRN 80-62-6 CMF C5 H8 O2

CM 15

CRN 79-10-7 CMF C3 H4 O2

CRN 60-24-2 CMF C2 H6 O S

HO-CH2-CH2-SH

CM 17

CRN 59-52-9 CMF C3 H8 O S2

 $\begin{array}{c} \text{SH} \\ \mid \\ \text{HS-CH}_2\text{-CH-CH}_2\text{-OH} \end{array}$ 

RN 365438-70-6 HCAPLUS
CN Pentanedioic acid, dihydrazide, polymer with 2,2bis(hydroxymethyl)butanoic acid, bis(isocyanatomethyl)cyclohexane,
1,3-dihydroxy-2-propanone, α-hydro-ω-hydroxypoly(oxy1,4-butanediyl), 2-mercaptoethanol and methyl 2-methyl-2propenoate, compd. with N,N-diethylethanamine (9CI) (CA INDEX NAME)

CM 1

CRN 121-44-8 CMF C6 H15 N

Et | Et-N-Et

CM 2

CRN 365438-69-3
CMF (C10 H14 N2 O2 . C6 H12 O4 . C5 H12 N4 O2 . C5 H8 O2 . (C4 H8 O) n H2 O . C3 H6 O3 . C2 H6 O S) x
CCI PMS

CM 3

CRN 42170-25-2 CMF C10 H14 N2 O2 CCI IDS

CRN 25190-06-1 CMF (C4 H8 O)n H2 O CCI PMS

HO 
$$\left[ (CH_2)_4 - O \right]_n$$
 H

CM 5

CRN 10097-02-6 CMF C6 H12 O4

CM 6

CRN 1508-67-4 CMF C5 H12 N4 O2

$$\begin{array}{c|c} & & & \circ & & \circ \\ \parallel & & & \parallel & & \parallel \\ \text{$\text{H}_2$N-NH-C- (CH$_2)} & 3 - \text{C-NH-NH}_2 \\ \end{array}$$

CM 7

CRN 96-26-4 CMF C3 H6 O3

CRN 80-62-6 CMF C5 H8 O2

CM 9

CRN 60-24-2 CMF C2 H6 O S

 $HO-CH_2-CH_2-SH$ 

RN 365438-72-8 HCAPLUS

CN Hexanedioic acid, polymer with 2-aminoethanethiol,
5-amino-1,3,3-trimethylcyclohexanemethanamine, butyl 2-propenoate,
N-(1,1-dimethyl-3-oxobutyl)-2-propenamide, 2,2-dimethyl-1,3propanediol, hexanedioic acid dihydrazide, 1,6-hexanediol,
2-hydroxyethyl 2-propenoate, 3-hydroxy-2-(hydroxymethyl)-2methylpropanoic acid, 5-isocyanato-1-(isocyanatomethyl)-1,3,3trimethylcyclohexane, methyl 2-methyl-2-propenoate,
2-propenenitrile and 2-propenoic acid, compd. with
N,N-diethylethanamine (9CI) (CA INDEX NAME)

CM 1

CRN 121-44-8 CMF C6 H15 N

CM 2

CRN 365438-71-7

CMF (C12 H18 N2 O2 . C10 H22 N2 . C9 H15 N O2 . C7 H12 O2 . C6 H14 N4 O2 . C6 H14 O2 . C6 H10 O4 . C5 H12 O2 . C5 H10 O4 . C5 H8 O3 . C5 H8 O2 . C3 H4 O2 . C3 H3 N . C2 H7 N S)x

CCI PMS

CM 3

CRN 4767-03-7 CMF C5 H10 O4

CM 4

CRN 4098-71-9 CMF C12 H18 N2 O2

CM 5

CRN 2873-97-4 CMF C9 H15 N O2

$$\begin{array}{c} {\rm O} \\ || \\ {\rm H_2C} = {\rm CH-C-NH} \qquad {\rm O} \\ | \qquad || \\ {\rm Me-C-CH_2-C-Me} \\ | \\ {\rm Me} \end{array}$$

CM 6

CRN 2855-13-2 CMF C10 H22 N2

$$\begin{array}{c} \text{Me} & \text{Me} \\ \text{Me} & \text{CH}_2 - \text{NH}_2 \\ \\ \text{NH}_2 & \text{NH}_2 \end{array}$$

CRN 1071-93-8 CMF C6 H14 N4 O2

CM 8

CRN 818-61-1 CMF C5 H8 O3

$$\begin{array}{c} {\rm O} \\ || \\ {\rm HO-CH_2-CH_2-O-C-CH} \end{array}$$

CM 9

CRN 629-11-8 CMF C6 H14 O2

$$^{\rm HO^-}$$
 (CH<sub>2</sub>)<sub>6</sub>-OH

CM 10

CRN 141-32-2 CMF C7 H12 O2

$$\begin{array}{c} \text{O} \\ \parallel \\ \text{n-BuO-C-CH} \end{array}$$

CM 11

CRN 126-30-7 CMF C5 H12 O2

CRN 124-04-9 CMF C6 H10 O4

 $HO_2C^-$  (CH<sub>2</sub>)<sub>4</sub> -  $CO_2H$ 

CM 13

CRN 107-13-1 CMF C3 H3 N

 $H_2C = CH - C = N$ 

CM 14

CRN 80-62-6 CMF C5 H8 O2

CM 15

CRN 79-10-7 CMF C3 H4 O2

о || но- с- сн== сн<sub>2</sub>

CM 16

CRN 60-23-1 CMF C2 H7 N S

 $H_2N-CH_2-CH_2-SH$ 

RN 365438-74-0 HCAPLUS

CN Hexanedioic acid, dihydrazide, polymer with 2-aminoethanethiol, 5-amino-1,3,3-trimethylcyclohexanemethanamine, 2,2-bis(hydroxymethyl)butanoic acid, butyl 2-propenoate, 1,6-diisocyanatohexane, N-(1,1-dimethyl-3-oxobutyl)-2-propenamide, α,α'-1,2-ethanediylbis[ω-hydroxypoly[oxy(1-oxo-

1,6-hexanediyl)]], methyl 2-methyl-2-propenoate and 2-propenoic acid, compd. with N,N-diethylethanamine (9CI) (CA INDEX NAME)

CM 1

CRN 121-44-8 CMF C6 H15 N

CM 2

CRN 365438-73-9
CMF (C10 H22 N2 . C9 H15 N O2 . C8 H12 N2 O2 . C7 H12 O2 . C6 H14 N4 O2 . C6 H12 O4 . (C6 H10 O2)n (C6 H10 O2)n C2 H6 O2 . C5 H8 O2 . C3 H4 O2 . C2 H7 N S)x
CCI PMS

CM 3

CRN 59692-54-5

CMF (C6 H10 O2)n (C6 H10 O2)n C2 H6 O2 CCI PMS

HO 
$$=$$
  $\begin{bmatrix} CH_2 \\ S \end{bmatrix}_5 - C = O = \begin{bmatrix} CH_2 \\ D \end{bmatrix}_n = CH_2 - CH_2 = \begin{bmatrix} O \\ CH_2 \\ CH_2 \end{bmatrix}_5 = \begin{bmatrix} O \\ CH_2 \\ D \end{bmatrix}_n = OH_2 - CH_2 = CH_2$ 

CM 4

CRN 10097-02-6 CMF C6 H12 O4

$$_{\rm CH_2-OH}^{\rm CH_2-OH}$$
 Et-C-CO<sub>2</sub>H  $_{\rm CH_2-OH}^{\rm CH_2-OH}$ 

CM 5

CRN 2873-97-4 CMF C9 H15 N O2

$$\begin{array}{c} {\rm O} \\ || \\ {\rm H_2C} = {\rm CH-C-NH} \qquad {\rm O} \\ || \\ {\rm Me-C-CH_2-C-Me} \\ || \\ {\rm Me} \end{array}$$

CRN 2855-13-2 CMF C10 H22 N2

$$\begin{array}{c} \text{Me} & \text{Me} \\ \text{Me} & \text{CH}_2 - \text{NH}_2 \\ \\ \text{NH}_2 & \\ \end{array}$$

CM 7

CRN 1071-93-8 CMF C6 H14 N4 O2

$$\begin{array}{c|c} & \circ & \circ \\ \parallel & \parallel \\ \text{$_{12}$N-NH-C-(CH$_{2})$_{4}-C-NH-NH$_{2}$} \end{array}$$

CM 8

CRN 822-06-0 CMF C8 H12 N2 O2

OCN-(CH<sub>2</sub>)<sub>6</sub>-NCO

CM 9

CRN 141-32-2 CMF C7 H12 O2

$$\begin{array}{c} \text{O} \\ \parallel \\ \text{n-BuO-C-CH-CH-CH}_2 \end{array}$$

CRN 80-62-6 CMF C5 H8 O2

$$\begin{array}{c|c} ^{H_2C} & \text{O} \\ \parallel & \parallel \\ \text{Me-} \text{C-} \text{C-} \text{OMe} \end{array}$$

CM 11

CRN 79-10-7 CMF C3 H4 O2

CM 12

CRN 60-23-1 CMF C2 H7 N S

 $H_2N-CH_2-CH_2-SH$ 

RN 365438-77-3 HCAPLUS

CN Butanoic acid, 2,2-bis(hydroxymethyl)-, polymer with 2-aminoethanethiol, 5-amino-1,3,3-trimethylcyclohexanemethanamine, butyl 2-propenoate, 1,6-diisocyanatohexane, N-(1,1-dimethyl-3-oxobutyl)-2-propenamide, α,α'-1,2-ethanediylbis[ω-hydroxypolyl[oxy(1-oxo-1,6-hexanediyl)]], methyl 2-methyl-2-propenoate and 2-propenoic acid, compd. with N,N-diethylethanamine (9CI) (CA INDEX NAME)

CM 1

CRN 121-44-8 CMF C6 H15 N

CM 2

CRN 365438-76-2 CMF (C10 H22 N2 . C9 H15 N O2 . C8 H12 N2 O2 . C7 H12 O2 . C6 H12 O4 . (C6 H10 O2)n (C6 H10 O2)n C2 H6 O2 . C5 H8 O2 . C3 H4 O2 . C2 H7 N S)x

CCI PMS

CM 3

CRN 59692-54-5

CMF (C6 H10 O2)n (C6 H10 O2)n C2 H6 O2

CCI PMS

HO — 
$$(CH_2)_5$$
 —  $(CH_2)_5$  —

CM 4

CRN 10097-02-6 CMF C6 H12 O4

$$_{\rm CH_2-OH}^{\rm CH_2-OH}$$
 Et-C-CO<sub>2</sub>H  $_{\rm CH_2-OH}^{\rm CH_2-OH}$ 

CM 5

CRN 2873-97-4 CMF C9 H15 N O2

$$\begin{array}{c} \text{O} \\ || \\ \text{H}_2\text{C} = \text{CH} - \text{C} - \text{NH} & \text{O} \\ | & || \\ \text{Me} - \text{C} - \text{CH}_2 - \text{C} - \text{Me} \\ | & \\ \text{Me} \end{array}$$

CM 6

CRN 2855-13-2 CMF C10 H22 N2

$$\begin{array}{c} \text{Me} & \text{Me} \\ \text{Me} & \text{CH}_2 - \text{NH}_2 \\ \\ \text{NH}_2 & \text{NH}_2 \end{array}$$

CRN 822-06-0 CMF C8 H12 N2 O2

OCN-(CH<sub>2</sub>)<sub>6</sub>-NCO

CM 8

CRN 141-32-2 CMF C7 H12 O2

$$\begin{array}{c}
O \\
\parallel \\
n-BuO-C-CH-CH-CH_2
\end{array}$$

CM 9

CRN 80-62-6 CMF C5 H8 O2

CM 10

CRN 79-10-7 CMF C3 H4 O2

CM 11

CRN 60-23-1

CMF C2 H7 N S

 $H_2N-CH_2-CH_2-SH$ 

RN 365438-79-5 HCAPLUS

CN Hexanedioic acid, polymer with 2,2-bis(hydroxymethyl)butanoic acid, butyl 2-propenoate, 1,6-diisocyanatohexane,
N-(1,1-dimethyl-3-oxobutyl)-2-propenamide, 2,2-dimethyl-1,3-propanediol, 2-hydroxyethyl 2-methyl-2-propenoate,
2-mercaptoethanol, methyl 2-methyl-2-propenoate,
3-methyl-1,5-pentanediol, pentanedioic acid dihydrazide and
2-propenoic acid, compd. with N,N-diethylethanamine (9CI) (CFINDEX NAME)

CM 1

CRN 121-44-8 CMF C6 H15 N

Et | | Et-N-Et

CM 2

CRN 365438-78-4
CMF (C9 H15 N O2 . C8 H12 N2 O2 . C7 H12 O2 . C6 H14 O2 . C6 H12 O4 . C6 H10 O4 . C6 H10 O3 . C5 H12 N4 O2 . C5 H12 O2 . C5 H8 O2 . C3 H4 O2 . C2 H6 O S)x
CCI PMS

CM 3

CRN 10097-02-6 CMF C6 H12 O4

 $_{\rm CH_2-OH}^{\rm CH_2-OH}$  Et-C-CO<sub>2</sub>H  $_{\rm CH_2-OH}^{\rm CH_2-OH}$ 

CM 4

CRN 4457-71-0 CMF C6 H14 O2

 $\begin{array}{c} & \text{Me} \\ | \\ \text{HO-} \ \text{CH}_2\text{--} \ \text{CH}_2\text{--} \ \text{CH}_2\text{--} \ \text{CH}_2\text{--} \ \text{OH} \end{array}$ 

CRN 2873-97-4 CMF C9 H15 N O2

$$\begin{array}{c} {\rm O} \\ || \\ {\rm H_2C} = {\rm CH-C-NH} \\ || \\ {\rm Me-C-CH_2-C-Me} \\ || \\ {\rm Me} \end{array}$$

CM 6

CRN 1508-67-4 CMF C5 H12 N4 O2

$$\begin{array}{c|c} & & & \circ & \\ \parallel & & \parallel & \\ \text{$_{\text{H}_2}$N-NH-C- (CH$_2)} & \text{$_{\text{3}}$-C-NH-NH$_2} \end{array}$$

CM 7

CRN 868-77-9 CMF C6 H10 O3

CM 8

CRN 822-06-0 CMF C8 H12 N2 O2

OCN-(CH<sub>2</sub>)<sub>6</sub>-NCO

CM 9

CRN 141-32-2 CMF C7 H12 O2

CRN 126-30-7 CMF C5 H12 O2

CM 11

CRN 124-04-9 CMF C6 H10 O4

$$HO_2C^-$$
 (CH<sub>2</sub>)<sub>4</sub> - CO<sub>2</sub>H

CM 12

CRN 80-62-6 CMF C5 H8 O2

$$\begin{array}{ccc} ^{H_2C} & \text{O} \\ & \parallel & \parallel \\ \text{Me-} & \text{C-} & \text{C-} & \text{OMe} \end{array}$$

CM 13

CRN 79-10-7 CMF C3 H4 O2

CM 14

CRN 60-24-2 CMF C2 H6 O S  $HO-CH_2-CH_2-SH$ 

CN

RN 365438-81-9 HCAPLUS

Pentanedioic acid, dihydrazide, polymer with 2,2-bis(hydroxymethyl)butanoic acid, bis(isocyanatomethyl)cyclohexane, 1,3-dihydroxy-2-propanone, α-hydro-ω-hydroxypoly(oxy-1,4-butanediyl), 2-hydroxyethyl 2-methyl-2-propenoate, 2-mercaptoethanol and methyl 2-methyl-2-propenoate, compd. with N,N-diethylethanamine (9CI) (CA INDEX NAME)

CM 1

CRN 121-44-8 CMF C6 H15 N

Et | Et- **N**- Et

CM 2

CRN 365438-80-8

CMF (C10 H14 N2 O2 . C6 H12 O4 . C6 H10 O3 . C5 H12 N4 O2 . C5 H8

O2 . (C4 H8 O)n H2 O . C3 H6 O3 . C2 H6 O S)x

CCI PMS

CM 3

CRN 42170-25-2 CMF C10 H14 N2 O2 CCI IDS



2 D1-CH2-NCO

CM 4

CRN 25190-06-1 CMF (C4 H8 O)n H2 O CCI PMS

HO 
$$\left[ - \left( CH_2 \right)_4 - O \right]_n$$
 H

CRN 10097-02-6 CMF C6 H12 O4

CM 6

CRN 1508-67-4 CMF C5 H12 N4 O2

$$\begin{array}{c|c} & & & \circ & \\ \parallel & & \parallel \\ \text{$_{12}$N-$NH-$C- (CH$_{2})$} & \text{$_{3}$-$C-$NH-$NH$_{2}$} \end{array}$$

CM 7

CRN 868-77-9 CMF C6 H10 O3

CM 8

CRN 96-26-4 CMF C3 H6 O3

$$0 \\ || \\ HO-CH_2-C-CH_2-OH$$

CM 9

CRN 80-62-6

CMF C5 H8 O2

 $^{\text{H}_2\text{C}}_{||}$   $^{\text{O}}_{||}$   $^{\text{Me}-\text{C}-\text{C}-\text{OMe}}$ 

CM 10

CRN 60-24-2 CMF C2 H6 O S

 $HO-CH_2-CH_2-SH$ 

RN 365438-83-1 HCAPLUS

CN Hexanedioic acid, polymer with 2-aminoethanethiol,
5-amino-1,3-trimethylcyclohexanemethanamine, butyl 2-propenoate,
N-(1,1-dimethyl-3-oxobutyl)-2-propenamide, 2,2-dimethyl-1,3propanediol, hexanedioic acid dihydrazide, 1,6-hexanediol,
2-hydroxyethyl 2-methyl-2-propenoate, 2-hydroxyethyl 2-propenoate,
3-hydroxy-2-(hydroxymethyl)-2-methylpropanoic acid,
5-isocyanato-1-(isocyanatomethyl)-1,3,3-trimethylcyclohexane,
methyl 2-methyl-2-propenoate, 2-propenenitrile and 2-propenoic
acid, compd. with N,N-diethylethanamine (9CI) (CA INDEX NAME)

CM 1

CRN 121-44-8 CMF C6 H15 N

Et | Et-N-Et

CM 2

CRN 365438-82-0
CMF (C12 H18 N2 O2 . C10 H22 N2 . C9 H15 N O2 . C7 H12 O2 . C6 H14 N4 O2 . C6 H14 O2 . C6 H10 O4 . C6 H10 O3 . C5 H12 O2 . C5 H10 O4 . C5 H8 O3 . C5 H8 O2 . C3 H4 O2 . C3 H3 N . C2 H7 N S)x

см з

CCI PMS

CRN 4767-03-7 CMF C5 H10 O4

CRN 4098-71-9 CMF C12 H18 N2 O2

CM 5

CRN 2873-97-4 CMF C9 H15 N O2

$$\begin{array}{c} \text{O} \\ || \\ \text{H}_2\text{C} = \text{CH} - \text{C} - \text{NH} & \text{O} \\ || \\ \text{Me} - \text{C} - \text{CH}_2 - \text{C} - \text{Me} \\ || \\ \text{Me} \end{array}$$

CM 6

CRN 2855-13-2 CMF C10 H22 N2

Me Me 
$$CH_2-NH_2$$
  $NH_2$ 

CM 7

CRN 1071-93-8

CMF C6 H14 N4 O2

CM 8

CRN 868-77-9 CMF C6 H10 O3

$$^{\rm H_2C}_{\parallel}$$
  $^{\rm O}_{\parallel}$   $^{\rm Me-}$   $^{\rm C-}$   $^{\rm C-}$   $^{\rm O-}$   $^{\rm CH_2-}$   $^{\rm CH_2-}$   $^{\rm OH}$ 

CM 9

CRN 818-61-1 CMF C5 H8 O3

$$\begin{array}{c} & \circ \\ || \\ \text{HO-CH}_2\text{--CH}_2\text{--}\text{--C-CH} \end{array} \text{CH}_2$$

CM 10

CRN 629-11-8 CMF C6 H14 O2

$$HO-(CH_2)_6-OH$$

CM 11

CRN 141-32-2 CMF C7 H12 O2

CM 12

CRN 126-30-7 CMF C5 H12 O2

$$\begin{array}{c} & \text{Me} \\ | \\ \text{HO-} \ \text{CH}_2 - \text{C-} \ \text{CH}_2 - \text{OH} \\ | \\ \text{Me} \end{array}$$

CRN 124-04-9 CMF C6 H10 O4

$$HO_2C^-$$
 (CH<sub>2</sub>)<sub>4</sub>-CO<sub>2</sub>H

CM 14

CRN 107-13-1 CMF C3 H3 N

$$H_2C = CH - C = N$$

CM 15

CRN 80-62-6 CMF C5 H8 O2

CM 16

CRN 79-10-7 CMF C3 H4 O2

CM 17

CRN 60-23-1 CMF C2 H7 N S  $H_2N-CH_2-CH_2-SH$ 

RN 365438-85-3 HCAPLUS
CN Hexanedioic acid, dihydrazide, polymer with 2-aminoethanethiol,
5-amino-1,3,3-trimethylcyclohexanemethanamine,
2,2-bis(hydroxymethyl)butanoic acid, butyl 2-propenoate,
1,6-diisocyanatohexane, N-(1,1-dimethyl-3-oxobutyl)-2-propenamide,
α,α'-1,2-ethanediylbis[ω-hydroxypoly[oxy(1-oxo1,6-hexanediyl)]], 2-hydroxyethyl 2-methyl-2-propenoate, methyl
2-methyl-2-propenoate and 2-propenoic acid, compd. with
N,N-diethylethanamine (9CI) (CA INDEX NAME)

CM 1

CRN 121-44-8 CMF C6 H15 N

CM 2

CRN 365438-84-2
CMF (C10 H22 N2 . C9 H15 N O2 . C8 H12 N2 O2 . C7 H12 O2 . C6 H14 N4 O2 . C6 H12 O4 . C6 H10 O3 . (C6 H10 O2)n (C6 H10 O2)n C2 H6 O2 . C5 H8 O2 . C3 H4 O2 . C2 H7 N S)x

CCI PMS

CM 3

CRN 59692-54-5 CMF (C6 H10 O2)n (C6 H10 O2)n C2 H6 O2 CCI PMS

CM 4

CRN 10097-02-6 CMF C6 H12 O4

$$CH_2-OH$$
  
 $|$   
 $Et-C-CO_2H$   
 $|$   
 $CH_2-OH$ 

CRN 2873-97-4 CMF C9 H15 N O2

$$\begin{array}{c} {\rm O} \\ || \\ {\rm H_2C} = {\rm CH-C-NH} \qquad {\rm O} \\ | \qquad || \\ {\rm Me-C-CH_2-C-Me} \\ | \qquad {\rm Me} \end{array}$$

CM 6

CRN 2855-13-2 CMF C10 H22 N2

$$\begin{array}{c|c} \text{Me} & \text{Me} \\ \text{Me} & \text{CH}_2 - \text{NH}_2 \\ \\ \text{NH}_2 & \\ \end{array}$$

CM 7

CRN 1071-93-8 CMF C6 H14 N4 O2

$$\begin{array}{c|c} & & & O & & O \\ \parallel & & & \parallel & & \parallel \\ H_2N-NH-C-(CH_2)_4-C-NH-NH_2 & & & & \end{array}$$

CM 8

CRN 868-77-9 CMF C6 H10 O3

CM 9

CRN 822-06-0 CMF C8 H12 N2 O2

OCN-(CH<sub>2</sub>)<sub>6</sub>-NCO

CM 10

CRN 141-32-2 CMF C7 H12 O2

CM 11

CRN 80-62-6 CMF C5 H8 O2

$$\begin{array}{ccc} ^{H_2C} & \text{O} \\ & \parallel & \parallel \\ \text{Me-} & \text{C--} & \text{C--} & \text{OMe} \end{array}$$

CM 12

CRN 79-10-7 CMF C3 H4 O2

CM 13

CRN 60-23-1 CMF C2 H7 N S

 $H_2N-CH_2-CH_2-SH$ 

RN 365438-88-6 HCAPLUS

CN Butanoic acid, 2,2-bis(hydroxymethyl)-, polymer with 2-aminoethanethiol, 5-amino-1,3,3-trimethylcyclohexanemethanamine, butyl 2-propenoate, 1,6-diisocyanatohexane, N-(1,1-dimethyl-3oxobutyl)-2-propenamide, α,α'-1,2ethanediylbis[ω-hydroxypoly[oxy(1-oxo-1,6-hexanediyl)]], 2-hydroxyethyl 2-methyl-2-propenoate, methyl 2-methyl-2-propenoate and 2-propenoic acid, compd. with N,N-diethylethanamine (9CI) (CA INDEX NAME)

CM 1

CRN 121-44-8 CMF C6 H15 N

CM 2

CRN 365438-87-5

CMF (C10 H22 N2 . C9 H15 N O2 . C8 H12 N2 O2 . C7 H12 O2 . C6 H12 O4 . C6 H10 O3 . (C6 H10 O2)n (C6 H10 O2)n C2 H6 O2 . C5 H8 O2 . C3 H4 O2 . C2 H7 N S)x

CCI PMS

CM 3

CRN 59692-54-5 CMF (C6 H10 O2)n (C6 H10 O2)n C2 H6 O2 CCI PMS

HO — 
$$\begin{bmatrix} & O & \\ & & \\ & & \\ & & \end{bmatrix}$$
 CH<sub>2</sub> — CH<sub>2</sub> —  $\begin{bmatrix} & O & \\ & & \\ & & \\ & & \end{bmatrix}$  OH

CM 4

CRN 10097-02-6 CMF C6 H12 O4

$$_{\rm CH_2-OH}^{\rm CH_2-OH}$$
  $_{\rm CH_2-OH}^{\rm CH_2-OH}$ 

CM 5

CRN 2873-97-4 CMF C9 H15 N O2

$$\begin{array}{c} {\rm O} \\ || \\ {\rm H_2C} = {\rm CH-C-NH} \\ | \\ | \\ {\rm Me-C-CH_2-C-Me} \\ | \\ {\rm Me} \end{array}$$

CRN 2855-13-2 CMF C10 H22 N2

$$\begin{array}{c} \text{Me} & \text{Me} \\ \text{Me} & \text{CH}_2 - \text{NH}_2 \\ \\ \text{NH}_2 & \text{NH}_2 \end{array}$$

CM 7

CRN 868-77-9 CMF C6 H10 O3

CM 8

CRN 822-06-0 CMF C8 H12 N2 O2

OCN-(CH<sub>2</sub>)<sub>6</sub>-NCO

CM 9

CRN 141-32-2 CMF C7 H12 O2

$$\begin{array}{c} \text{O} \\ \parallel \\ \text{n-BuO-C-CH------} \text{CH}_2 \end{array}$$

CRN 80-62-6 CMF C5 H8 O2

$$^{\text{H}_2\text{C}}_{\parallel}$$
  $^{\text{O}}_{\parallel}$   $^{\text{Me}-\text{C}-\text{C}-\text{OMe}}$ 

CM 11

CRN 79-10-7 CMF C3 H4 O2

CM 12

CRN 60-23-1 CMF C2 H7 N S

 $H_2N-CH_2-CH_2-SH$ 

IC ICM C08F283-00

ICS C08F002-44; C08G018-28; C08L075-04; C09D151-08

CC 42-7 (Coatings, Inks, and Related Products)

IT 60-24-2DP, acrylic polyurethanes 80-62-6DP, Methyl methacrylate, acrylic polyurethanes 96-26-4DP, Bis(hydroxymethyl)ketone, acrylic polyurethanes 101-43-9DP, Cyclohexyl methacrylate, acrylic polyurethanes 629-11-8DP, 1,6-Hexanediol, reaction products with dimer acids, acrylic polyurethanes 868-77-9DP, 2-Hydroxyethyl methacrylate, acrylic polyurethanes 2873-97-4DP. Diacetoneacrylamide, acrylic polyurethanes 5124-30-1DP, Hydrogenated MDI, acrylic polyurethanes 10097-02-6DP, 2,2-Dimethylolbutyric acid, acrylic polyurethanes 365438-66-0P 365438-68-2P 365438-70-6P 365438-72-8P 365438-74-0P 365438-77-3P

365438-79-5P 365438-81-9P 365438-83-1P

365438-85-3P 365438-88-6P

(manufacture of aqueous acrylic urethane polymer emulsions without using organic solvents for haze-free flexible coatings)

L21 ANSWER 10 OF 25 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER:

1993:549531 HCAPLUS

DOCUMENT NUMBER:

119:149531

TITLE:

Photosensitive compositions useful

for making presensitized lithographic plates

INVENTOR (S):

Nakai, Hideyuki; Suzuki, Toshitsugu; Matsumura, Tomoyuki

PATENT ASSIGNEE(S): SOURCE:

Konishiroku Photo Ind, Japan Jpn. Kokai Tokkyo Koho, 12 pp.

CODEN: JKXXAF

DOCUMENT TYPE: LANGUAGE:

Patent Japanese

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 04362647	A2	19921215	JP 1991-164992	
				1991
				0610
PRIORITY APPLN. INFO.:			JP 1991-164992	
				1991
				0610

AB In the title composition comprising a compound which generates acid upon irradiation with actinic rays and a compound having a bond decomposable by the acid, the compound which generates acid upon irradiation with actinic rays is a condensate of o-naphthoquinonediazidosulfonic acid halide and an oxime-containing compound The compns. show good stability in photosensitivity after exposure and underdevelopability (developability with respect to developers with reduced developing capacity).

IT 149671-27-2P

> (preparation of, photosensitive composition containing, for lithog. plate making)

RN149671-27-2 HCAPLUS

CN Formic acid, methyl ester, polymer with 3-mercapto-1,2-propanediol (9CI) (CA INDEX NAME)

CM 1

CRN 107-31-3 CMF C2 H4 O2

O CH- O- CH3

CM 2

CRN 96-27-5 CMF C3 H8 O2,

OH HS-CH2-CH-CH

IC ICM G03F007-022

CC 74-6 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)

ST photosensitive compn naphthoquinonediazide compd; oxime condensate photosensitive compn; presensitized lithog

plate photosensitive compn

IT Lithographic plates

(presensitized, photosensitive composition for, containing

naphthoquinonediazide acid-generator)

IT 146905-39-7P 146905-40-0P 146905-41-1P

(preparation of, photosensitive composition containing,

acid-generator, for lithog. plate making)

IT 69468-60-6P 78537-86-7P 115324-80-6P 115815-82-2P

116745-41-6P 117646-94-3P 117647-26-4P 117647-27-5P 117992-19-5P 118188-70-8P 119201-95-5P 149671-27-2P

149671-30-7P

(preparation of, photosensitive composition containing, for lithog. plate making)

L21 ANSWER 11 OF 25 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1993:244642 HCAPLUS

DOCUMENT NUMBER: 118:244642

TITLE: Photosensitive compositions useful

for making presensitized lithographic plates

INVENTOR(S):

Nakai, Hideyuki; Akyama, Takeo Konishiroku Photo Ind, Japan Inn Kokai Tokkyo Kobo, 11 pp

PATENT ASSIGNEE(S): SOURCE:

Jpn. Kokai Tokkyo Koho, 11 pp. CODEN: JKXXAF

DOCUMENT TYPE:

LANGUAGE:

Patent Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
TD 04050540				
JP 04362643	A2	19921215	JP 1991-164993	
				1991
				0610
PRIORITY APPLN. INFO.:			JP 1991-164993	
				1991
				0610

OTHER SOURCE(S):

MARPAT 118:244642

GΙ

AB The title compns. comprise a compound I (A = Q, Q1, Q2; X = halo; k, l, m = 1-3; B = monovalent anion; R1 = H, alkyl, aryl,

```
aralkyl; aromatic heterocycle) generating an acid by irradiation with
     active rays and another having a bond which is decomposed
     by the acid. The compns. show good
     photosensitivity and under-developability.
IT
     115324-68-0P
        (preparation of, photosensitive composition containing, for lithog.
        plate making)
RN
     115324-68-0 HCAPLUS
CN
     1,2-Propanediol, 3-mercapto-, polymer with trimethoxymethane (9CI)
       (CA INDEX NAME)
     CM
          1
     CRN
         149-73-5
     CMF C4 H10 O3
     OMe
MeO-CH-OMe
     CM
     CRN 96-27-5
     CMF C3 H8 O2 S
        OH
HS-CH2-CH-CH2-OH
IC
     ICM G03F007-004
     ICS G03F007-029; G03F007-039; H01L021-027
     74-6 (Radiation Chemistry, Photochemistry, and
CC
     Photographic and Other Reprographic Processes)
ST
     photosensitive compn acid generating compd;
     presensitized lithog plate photosensitive compn
TT
     Lithographic plates
        (presensitized, photosensitive composition for, containing
        acid-generator)
IT
     142515-37-5
                   147360-65-4
                                 147360-66-5
                                               147769-57-1
        (photosensitive composition containing, acid-generator, for
        lithog. plate making)
IT
     69468-60-6P
                   78537-86-7P
                                 96758-30-4P
                                               115157-62-5P
     115324-68-0P
                    115324-80-6P
                                   116745-41-6P
                                                  117646-94-3P
     117647-26-4P
                    117647-27-5P
                                   117992-19-5P
        (preparation of, photosensitive composition containing, for lithog.
        plate making)
L21 ANSWER 12 OF 25 HCAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                         1993:244641 HCAPLUS
DOCUMENT NUMBER:
                         118:244641
TITLE:
                         Photosensitive compositions useful
                         for making presensitized lithographic plates
INVENTOR(S):
                         Nakai, Hideyuki
                         Konishiroku Photo Ind, Japan
PATENT ASSIGNEE(S):
SOURCE:
                         Jpn. Kokai Tokkyo Koho, 9 pp.
```

CODEN: JKXXAF

DOCUMENT TYPE:

Patent

LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 04362642	A2	19921215	JP 1991-164991	
				1991
				0610
PRIORITY APPLN. INFO.:			JP 1991-164991	
				1991
				0610

OTHER SOURCE(S): MARPAT 118:244641

For diagram(s), see printed CA Issue.

The title compns. comprise a compound which generates an acid by irradiation with active rays, a compound having ≥1 bond which is decomposed by the acid, and a nitro compound I [A = (substituted) aromatic or heterocyclic ring having 5-14 ring-forming atoms; R1-3 = H, (substituted) alkyl, aryl, aralkyl] forming an amine compound by irradiation with active rays. The compns. show good stability in photosensitivity after exposure and dot-reproducibility.

IT 115324-68-0P

> (preparation of, photosensitive composition containing, for lithog. plate making)

115324-68-0 HCAPLUS RN

1,2-Propanediol, 3-mercapto-, polymer with trimethoxymethane (9CI) CN (CA INDEX NAME)

CM 1

CRN 149-73-5 CMF C4 H10 O3

OMe MeO-CH-OMe

> CM 2

CRN 96-27-5 CMF C3 H8 O2 S

OH  $HS-CH_2-CH-CH_2-OH$ 

ICM G03F007-004 IC

ICS G03F007-004; G03F007-029; G03F007-039

CC 74-6 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)

```
photosensitive compn amine forming compd; nitro compd
ST
     photosensitive compn; presensitized lithog plate
     photosensitive compn
TΤ
     Lithographic plates
        (presensitized, photosensitive composition for, containing
        amine-generator and acid-generator)
IT
     69432-40-2
                 72015-26-0
                               93641-24-8
        (photosensitive composition containing, acid-generator, for
        lithog. plate making)
IT
     74109-34-5
                74109-35-6
                               119137-03-0 147382-82-9
        (photosensitive composition containing, amine-generator, for
        lithog. plate making)
IT
     69468-60-6P
                  78537-86-7P
                                96758-30-4P 115157-62-5P
     115324-68-0P
                  115324-80-6P
                                 116745-41-6P 117646-94-3P,
     1,1-Dimethoxycyclohexane-triethylene glycol copolymer
     117647-26-4P, 1,1-Dimethoxycyclohexane-triethylene glycol
     copolymer, sru 117647-27-5P 117992-19-5P
        (preparation of, photosensitive composition containing, for lithog.
        plate making)
L21 ANSWER 13 OF 25 HCAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                        1993:222963 HCAPLUS
DOCUMENT NUMBER:
                        118:222963
TITLE:
                        Photosensitive compositions useful
                        for making presensitized lithographic plates
INVENTOR(S):
                        Nakai, Hideyuki; Akyama, Takeo
PATENT ASSIGNEE(S):
                        Konishiroku Photo Ind, Japan
SOURCE:
                        Jpn. Kokai Tokkyo Koho, 10 pp.
                        CODEN: JKXXAF
DOCUMENT TYPE:
                        Patent
LANGUAGE:
                        Japanese
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
     PATENT NO.
                        KIND
                               DATE
                                          APPLICATION NO.
                                                                  DATE
     -----
     ------
     JP 04362644
                        A2
                               19921215
                                           JP 1991-164994
                                                                  1991
                                                                  0610
PRIORITY APPLN. INFO.:
                                           JP 1991-164994
                                                                  1991
                                                                  0610
OTHER SOURCE(S):
                        MARPAT 118:222963
    The title compns. comprise 2-halomethyl-1,3,4-oxadiazole
     compound having a (substituted) naphthofuryl group at its 5 position
     directly or through vinyl group which generating an acid by
     irradiation with active rays and a compound having a bond which is
     decomposed by the acid. The compns.
    provide presensitized lithog. plates with high photosensitivity.
     115324-68-0P, Methyl orthoformate-thioglycerol copolymer
TT
        (preparation of, photosensitive composition containing, for lithog.
       plate making)
RN
     115324-68-0 HCAPLUS
     1,2-Propanediol, 3-mercapto-, polymer with trimethoxymethane (9CI)
CN
       (CA INDEX NAME)
```

CRN 149-73-5 CMF C4 H10 O3

OMe | MeO-CH-OMe

CM 2

CRN 96-27-5 CMF C3 H8 O2 S

ОН | HS- CH<sub>2</sub>- CH- CH<sub>2</sub>- ОН

IC ICM G03F007-004

ICS G03F007-029; G03F007-039

CC 74-6 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)
ST photosensitive compn halomethyloxadiazole deriv; acid decomposable compd photosensitive compn; presensitized lithog plate photosensitive compn

IT Lithographic plates

(presensitized, photosensitive composition for, containing oxadiazole acid-generator)

IT 147382-83-0 147382-84-1 147382-85-2 147382-86-3 (photosensitive composition containing, acid-generator, for lithog. plate making)

TT 69468-60-6P, Benzaldehyde-triethylene glycol copolymer 78537-86-7P, 1,2,6-Hexanetriol-methyl orthoformate copolymer 96758-30-4P 115157-62-5P, 1,2,6-Hexanetriol-tetramethyl orthocarbonate copolymer 115324-68-0P, Methyl orthoformate-thioglycerol copolymer 115324-80-6P, Methyl orthoformate-thioglycerol copolymer, sru 116745-41-6P, 1,2,6-Hexanetriol-methyl orthoformate copolymer, sru 117646-94-3P, 1,1-Dimethoxycyclohexane-triethylene glycol copolymer 117647-26-4P 117647-27-5P, Benzaldehyde-triethylene glycol copolymer, sru 117992-19-5P, 1,2,6-Hexanetrioltetramethyl orthocarbonate copolymer, sru (preparation of, photosensitive composition containing, for lithog. plate making)

L21 ANSWER 14 OF 25 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER:

1989:174024 HCAPLUS

DOCUMENT NUMBER:

110:174024

TITLE:

Preparation of acid-

cleavable epoxide addition polymers

containing acetal linkages

INVENTOR(S):

Klee, Joachim; Hoerhold, Hans Heinrich; Hopf,

Gert; Rode, Peter

PATENT ASSIGNEE(S):

Friedrich-Schiller-Universitaet, Ger. Dem.

Rep.

SOURCE:

Ger. (East), 4 pp.

CODEN: GEXXA8

DOCUMENT TYPE:

Patent

LANGUAGE:

German

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
DD 257435	A1	19880615	DD 1987-299685	
23/433	AI	19000013	DD 1967-299065	1987
				0204
PRIORITY APPLN. INFO.:			DD 1987-299685	
				1987
				0204

AB The title polymers, useful as adhesives, passivation materials in the elec. industry, resists, etc., are prepared by the reaction of diepoxides with acetal compds. such as RCH(OCH2CH2NHR1)2 or RCH(OCH2CH2SH)2 (R = H, Me, vinyl, Ph, etc.; R1 = Ph, PhCH2, Me, etc.). Heating 14.69 mmol bisphenol A diglycidyl ether and 14.69 mmol 1,7-bis(benzylamino)-3,5dioxaheptane 24 h at 60° gave a polymer (mol. weight 11,500) which was soluble in THF, pyridine, etc., and cleavable by alc. HCl.

IT 119889-66-6P

(preparation of acid-cleavable)

RN 119889-66-6 HCAPLUS

CN Ethanethiol, 2,2'-[methylenebis(oxy)]bis-, polymer with 2,2'-[(1-methylethylidene)bis(4,1-phenyleneoxymethylene)]bis[oxira ne] (9CI) (CA INDEX NAME)

CM 1

CRN 16128-67-9 CMF C5 H12 O2 S2

 $HS-CH_2-CH_2-O-CH_2-O-CH_2-CH_2-SH$ 

CM 2

CRN 1675-54-3 CMF C21 H24 O4

IC ICM C08G059-10

35-8 (Chemistry of Synthetic High Polymers)

Section cross-reference(s): 37

ST epoxy acetal resin acid cleavage; depolymn acid epoxy acetal resin; polymn addn epoxy acetal depolymn; resist epoxy acetal resin depolymn; adhesive epoxy acetal
resin depolymn; passivation epoxy acetal resin depolymn

IT Passivation

(acetal compound-diepoxide addition products for, preparation of acid-cleavable)

IT Adhesives

Resists

(acetal compound-diepoxide addition products, preparation of acid-cleavable)

IT Acetals

(diamino and dithiol derivs., addition products with diepoxides, preparation of acid-cleavable)

IT Epoxy resins, compounds

(reaction products, with diamino and dithiol derivs. of acetals, preparation of acid-cleavable)

IT 119889-66-6P 119923-62-5P 120127-26-6P (preparation of acid-cleavable)

L21 ANSWER 15 OF 25 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1989:125479 HCAPLUS

DOCUMENT NUMBER: 110:125479

TITLE: Photosensitive compositions, and

lithographic plate containing the same

INVENTOR(S): Nakai, Hideyuki; Sasa, Nobumasa; Matsubara,

Shinichi; Urano, Toshiyoshi; Murakami, Sachiko

PATENT ASSIGNEE(S): Konica Co., Japan; Mitsubishi Chemical

Industries Co., Ltd.

SOURCE: Jpn. Kokai Tokkyo Koho, 13 pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 63097944	A2	19880428	JP 1986-243861	
	•			1986
				1014
PRIORITY APPLN. INFO.:			JP 1986-243861	
				1986
				1014

AB The title composition contains (a) a compound that liberates acid on irradiation with activating radiation, (b) a compound having acid-cleavable bond(s), (c) novolak resin, and (d) a polymer having structural unit -CR1R2CR3 (CONR4XnYOH) - [R1-R2 = H, alkyl, carboxy; R3 = H, halo, alkyl; R4 = H, alkyl, Ph, aralkyl; Y = (substituted) phenylene, naphthylene; X = connecting group between N and aromatic C atoms; n = 0-5]. Lithog. plates having photosensitive layer containing the above composition is also claimed. The photosensitive composition and the lithog. plates have especially wide range of tolerance of development conditions. Thus, a solution containing acid-cleavable 1:1 dimethoxycyclohexane-triethylene glycol copolymer 2.14, novolak from phenol 2, m-cresol 4.8, and p-cresol 3.2 and HCHO 1.91, and 3:3:4 acrylonitrile-p-hydroxymethacrylanilide-Me methacrylate copolymer 3.83, Victoria Pure Blue BOH 0.05, and 2-trichloromethyl-5-[β-(2-benzofuryl)vinyl-1,3,4-oxadiazole

```
0.027 g was coated on an Al plate to obtain a lithog. plate. This
     plate was sensitometrically exposed to UV, and developed using
     alkaline developers whose dilution factor of stock solution ranged from 3 to
     15. Lithog. plates good for use were obtained with any of these
     developers, using same development time and temperature High
     resistance to plate cleaner and developing ink were also
     confirmed.
IT
     115324-68-0
        (photosensitive lithog. plates containing, acid-
        cleavable)
     115324-68-0 HCAPLUS
RN
     1,2-Propanediol, 3-mercapto-, polymer with trimethoxymethane (9CI)
CN
       (CA INDEX NAME)
     CM
          1
     CRN 149-73-5
     CMF C4 H10 O3
    OMe
MeO-CH-OMe
     CM
          2
     CRN 96-27-5
     CMF C3 H8 O2 S
        OH
HS-CH_2-CH-CH_2-OH
IC
     ICM G03C001-72
     ICS G03C001-72; G03F007-02
CC
     74-6 (Radiation Chemistry, Photochemistry, and
     Photographic and Other Reprographic Processes)
     Section cross-reference(s): 38
     lithog plate development condition tolerance; photosensitive
ST
     compn acid generating compd
IT
     Lithographic plates
        (photosensitive compns., with wide tolerance to
        development conditions)
IT
     Siloxanes and Silicones, uses and miscellaneous
        (photosensitive lithog. plates containing, acid-
        cleavable)
IT
     117646-96-5
        (photosensitive lithog. plates containing acid-
        cleavable compound and, for acid generation by activating
        radiations)
IT
     69468-60-6
                  78537-86-7
                               115157-62-5 115324-68-0
     117646-94-3
        (photosensitive lithog. plates containing, acid-
        cleavable)
L21 ANSWER 16 OF 25 HCAPLUS COPYRIGHT 2006 ACS on STN
```

ACCESSION NUMBER:

1989:105034 HCAPLUS

DOCUMENT NUMBER:

110:105034

TITLE:

Photosensitive compositions, and

lithographic plates containing the same

Nakai, Hideyuki; Matsubara, Shinichi; Urano,

Toshiyoshi; Murakami, Sachiko

PATENT ASSIGNEE(S):

Konica Co., Japan; Mitsubishi Chemical

Industries Co., Ltd.

SOURCE:

Jpn. Kokai Tokkyo Koho, 8 pp.

CODEN: JKXXAF

DOCUMENT TYPE: LANGUAGE:

INVENTOR (S):

Patent Japanese

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 63149638	A2	19880622	JP 1986-297305	
				1986
				1212
PRIORITY APPLN. INFO.:			JP 1986-297305	
				1986
				1212

Compds. that liberate acids and those that liberate amines by AΒ irradiation with activating radiation, and compds. that have a bond or bonds cleavable by the acids, are contained in the title compns. The title lithog. plates have a photosensitive layer or layers containing these compns. These compns. maintain a stable sensitivity after the exposure, and have high reproducibility of halftone dots. an acid-cleavable polymeric compound with units of the formula -(CH2CH2O)3(1,1-cyclohexylene)0-2.14, a cresol novolak 5.74, 2-trichloromethyl-5-[β-(2-benzofuryl)vinyl]-1,3,4-oxadiazole (acid-liberating compound) 0.27, Victoria Pure Blue BOH 0.05, and acetanilide (amine-liberating compound) 0.05 g dissolved in Me cellosolve were applied on anodized Al plates to obtain photosensitive lithog. plates. These plates showed no change of sensitivity upon a 90-min postexposure standing, either before or after a 2-day preexposure storage of the plates at 40° and 80% relative humidity.

IT 115324-68-0

> (photosensitive lithog. plates containing acid-generating compds. and amine generators and, for stable sensitivity)

RN115324-68-0 HCAPLUS

1,2-Propanediol, 3-mercapto-, polymer with trimethoxymethane (9CI) CN (CA INDEX NAME)

CM 1

CRN 149-73-5 CMF C4 H10 O3

OMe MeO-CH-OMe CM

2

```
CRN 96-27-5
     CMF C3 H8 O2 S
        OH
HS-CH_2-CH-CH_2-OH
IC
     ICM G03C001-72
     ICS G03C001-00
     74-6 (Radiation Chemistry, Photochemistry, and
CC
     Photographic and Other Reprographic Processes)
IT
     Lithographic plates
        (photosensitive compns. containing acid-generators and
        acid-cleavable compds. and amine generators
        for, for stable photosensitivity)
ΙT
     112-35-6, Triethylene glycol monomethyl ether
                                                      122-99-6, Phenyl
     cellosolve
        (condensation of, with acetal, acid-cleavable
        compds. for photosensitive lithog. plates from)
IT
     931-94-2, 1,1-Dimethoxycyclopentane
                                            933-40-4,
     1,1-Dimethoxycyclohexane
        (condensation of, with glycol derivative, acid-
        cleavable compds. for photosensitive lithog. plates
        from)
IT
     93641-24-8
        (photosensitive lithog. plates containing acid-
        cleavable compds. and amine-generating compds. and)
IT
     103-84-4, Acetanilide
        (photosensitive lithog. plates containing acid-generating compds.
        acid-cleavable compds. and, for stable
        sensitivity)
IT
     620-71-3, Propionanilide
        (photosensitive lithog. plates containing acid-generating compds.
        and acid-cleavable compds. and, for stable
        sensitivity)
IT
     69468-60-6
                  78537-86-7
                               96758-30-4 115324-68-0
     115324-80-6
                   115815-82-2
                                 116745-41-6
                                                117646-94-3
     117647-26-4
                   117647-27-5
                                 117992-19-5
                                                118188-70-8
     119201-95-5
        (photosensitive lithog. plates containing acid-generating compds.
        and amine generators and, for stable sensitivity)
L21 ANSWER 17 OF 25 HCAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                         1989:31459 HCAPLUS
DOCUMENT NUMBER:
                         110:31459
TITLE:
                         Photosensitive compositions, and
                         lithographic plates containing the same
                         Nakai, Hideyuki; Matsubara, Shinichi; Urano,
INVENTOR(S):
                         Toshiyoshi; Murakami, Sachiko
PATENT ASSIGNEE(S):
                         Konica Co., Japan; Mitsubishi Chemical
                         Industries Co., Ltd.
SOURCE:
                         Jpn. Kokai Tokkyo Koho, 9 pp.
                         CODEN: JKXXAF
DOCUMENT TYPE:
                         Patent
LANGUAGE:
                         Japanese
```

FAMILY ACC. NUM. COUNT: 1 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 63149640	A2	19880622	JP 1986-297307	
				1986
				1212
PRIORITY APPLN. INFO.:			JP 1986-297307	
				1986
				1212

AB Compds. that liberate acids, amines that are not decomposed by irradiation with activating radiations and scavenge the acids, and compds. that have bond(s) cleavable by the acids, are contained in the title compns. The title lithog. plates have photosensitive layer(s) containing these compns. These compns. provide sensitivity maintained stably after the exposure, and high reproducibility of halftone dots.

IT 115324-68-0P

(photosensitive lithog. plates containing acid-generating compds., amines and, for stable sensitivity, preparation of)

RN 115324-68-0 HCAPLUS

CM 1

CRN 149-73-5 CMF C4 H10 O3

OMe | MeO-CH-OMe

CM 2

CRN 96-27-5 CMF C3 H8 O2 S

 $\begin{array}{c} \text{OH} \\ | \\ \text{HS-CH}_2\text{-CH-CH}_2\text{-OH} \end{array}$ 

IC ICM G03C001-72 ICS G03C001-00

CC 74-6 (Radiation Chemistry, Photochemistry, and **Photographic** and Other Reprographic Processes)

IT Lithographic plates

(photosensitive compns., acid-generators, acid-cleavable compds. and amines contained in, for stable photosensitivity)

IT 112-35-6, Triethylene glycol monomethyl ether 122-99-6, Phenyl cellosolve

(condensation of, with acetal, acid-cleavable compds. for photosensitive lithog. plates from)

IT 931-94-2, 1,1-Dimethoxycyclopentane 933-40-4,

1,1-Dimethoxycyclohexane

(condensation of, with glycol derivative, acidcleavable compds. for photosensitive lithog. plates from)

IT 93641-24-8

(photosensitive lithog. plates containing acidcleavable compds., amines and)

IT 106-50-3, p-Phenylenediamine, uses and miscellaneous 150-13-0, p-Aminobenzoic acid 1122-58-3, 4-Dimethylaminopyridine

(photosensitive lithog. plates containing acid-generating compds. acid-cleavable compds. and, for stable

sensitivity)

IT 69468-60-6P 78537-86-7P 96758-30-4P 115157-62-5P 115324-68-0P 115324-80-6P 115815-82-2P 116745-41-6P 117646-94-3P 117647-26-4P 117647-27-5P 117992-19-5P

118188-70-8P

(photosensitive lithog. plates containing acid-generating compds., amines and, for stable sensitivity, preparation of)

L21 ANSWER 18 OF 25 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER:

1989:31458 HCAPLUS

DOCUMENT NUMBER:

110:31458

TITLE:

Photosensitive compositions, and

lithographic plates containing the same

INVENTOR(S): Nakai, Hideyuki; Matsubara, Shinichi; Urano,

Toshiyoshi; Murakami, Sachiko

PATENT ASSIGNEE(S):

Konica Co., Japan; Mitsubishi Chemical

Industries Co., Ltd.

SOURCE:

Jpn. Kokai Tokkyo Koho, 10 pp.

CODEN: JKXXAF

DOCUMENT TYPE:

Patent Japanese

LANGUAGE:

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 63149639	A2	19880622	JP 1986-297306	
				1986
				1212
PRIORITY APPLN. INFO.:			JP 1986-297306	
				1986
				1212

GΙ

AB Compds. that liberate acids, those that are not decomposed by irradiation with activating radiations and liberate amines by hydrolysis, and compds. that have bond(s) cleavable by

```
the acids, are contained in the title compns.
     The title lithog. plates have photosensitive layer(s) containing these
     compns. These compns. provide sensitivity
     maintained stably after the exposure, and high reproducibility of
     halftone dots.
IT
     115324-68-0P
        (photosensitive lithog, plates containing acid-generating compds.,
        amine generators and, for stable sensitivity, preparation of)
RN
     115324-68-0 HCAPLUS
CN
     1,2-Propanediol, 3-mercapto-, polymer with trimethoxymethane (9CI)
       (CA INDEX NAME)
     CM
     CRN
         149-73-5
     CMF C4 H10 O3
    OMe
MeO-CH-OMe
          2
     CM
         96-27-5
     CRN
     CMF C3 H8 O2 S
        OH
HS-CH_2-CH-CH_2-OH
IC
     ICM G03C001-72
     ICS G03C001-00
CC
     74-6 (Radiation Chemistry, Photochemistry, and
     Photographic and Other Reprographic Processes)
IT
     Lithographic plates
        (photosensitive compns., acid-generators,
        acid-cleavable compds. and hydrolyzable amine
        generators contained in, for stable photosensitivity)
IT
     112-35-6, Triethylene glycol monomethyl ether
                                                     122-99-6, Phenyl
        (condensation of, with acetal, acid-cleavable
        compds. for photosensitive lithog. plates from)
IT
     931-94-2, 1,1-Dimethoxycyclopentane
                                           933-40-4,
     1,1-Dimethoxycyclohexane
        (condensation of, with glycol derivative, acid-
        cleavable compds. for photosensitive lithog. plates
        from)
TΤ
     93641-24-8
        (photosensitive lithog. plates containing acid-
        cleavable compds., amine-generating compds. and)
IT
     783-08-4
        (photosensitive lithog. plates containing acid-generating compds.
        acid-cleavable compds. and, for stable
        sensitivity)
IT
     69468-60-6P
                   78537-86-7P
                                 96758-30-4P
                                               115157-62-5P
```

115324-68-0P 115324-80-6P 115815-82-2P 116745-41-6P 117646-94-3P 117647-26-4P 117647-27-5P 117992-19-5P

118188-70-8P

(photosensitive lithog. plates containing acid-generating compds., amine generators and, for stable sensitivity, preparation of)

L21 ANSWER 19 OF 25 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER:

1988:560638 HCAPLUS

DOCUMENT NUMBER:

109:160638

TITLE:

Photosensitive composition and

INVENTOR (S):

positive-type presensitized lithographic plate

Urano, Toshiyoshi; Murakami, Sachiko; Nakai,

Hideyuki; Sasa, Nobumasa

PATENT ASSIGNEE(S):

Mitsubishi Chemical Industries Co., Ltd.,

Japan; Konica Co.

SOURCE:

Jpn. Kokai Tokkyo Koho, 12 pp.

CODEN: JKXXAF

DOCUMENT TYPE:

Patent

LANGUAGE:

Japanese

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 63056650	A2	19880311	JP 1986-200006	
				1986
				0828
PRIORITY APPLN. INFO.:			JP 1986-200006	
				1986
				0828

GI

- AB The title photosensitive composition contains a polymer having a structural unit of I [R1, R2 = H, alkyl, carboxylic acid; R3 = H, halogen, alkyl; R4 = H, alkyl, Ph, aralkyl; Y = phenylene or naphthylene which may have a substituent; X = divalent organic group; n = 0-5]. A presensitized lithog. plate has a photosensitive layer consisting of the polymer, a compound which produces an acid by light exposure, and a compound (having ≥1 bond which increases solubility relative to an alkali solution) capable of being decomposed by acids. A highly sensitive and highly chemical resistant pos.-type photosensitive composition and a presensitized lithog. plate therefrom are obtained.
- IT 115324-68-0P

(preparation of, pos. presensitized lithog. plate from)

```
RN
     115324-68-0 HCAPLUS
     1,2-Propanediol, 3-mercapto-, polymer with trimethoxymethane (9CI)
CN
       (CA INDEX NAME)
     CM
     CRN
         149-73-5
     CMF C4 H10 O3
     OMe
MeO-CH-OMe
     CM
          2
     CRN
         96-27-5
     CMF C3 H8 O2 S
        OH
HS-CH_2-CH-CH_2-OH
IC
     ICM G03C001-72
     ICS G03F007-08
CC
     74-6 (Radiation Chemistry, Photochemistry, and
     Photographic and Other Reprographic Processes)
ST
     photosensitive compn; pos presensitized lithog plate
IT
     Lithographic plates
        (presensitized, pos., composition for, for improving
        sensitivity and chemical resistance)
IT
     19243-95-9P
                   27931-11-9P, N-(5-Hydroxy-α-naphthyl) methacryl
     amide
        (preparation and reaction of, pos.-type photosensitive compn
        . from, for presensitized lithog. plate)
IT
     78537-86-7P 115324-68-0P 116694-71-4P
        (preparation of, pos. presensitized lithog. plate from)
     75-78-5, Dichlorodimethyl silane 80-62-6, Methyl methacrylate
IT
     83-55-6, 5-Amino-\alpha-naphthol 96-27-5, Thioglycerol
     104-15-4, p-Toluene sulfonic acid, reactions 106-69-4,
     1,2,6-Hexanetriol
                         112-60-7, Tetraethylene glycol 123-30-8,
     p-Hydroxy aniline
                         140-88-5
                                    149-73-5, Trimethyl orthoformate
     150-76-5, Hydroquinone monomethyl ether
                                              589-29-7, p-Xylene
             868-77-9, 2-Hydroxyethyl methacrylate
                                                     920-46-7,
     Methacrylic acid chloride
        (reaction of, pos.-type photosensitive composition from,
        for presensitized lithog. plate)
L21 ANSWER 20 OF 25 HCAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                         1988:464377 HCAPLUS
DOCUMENT NUMBER:
                         109:64377
TITLE:
                         Color photoimaging composition for
                         preparing color proof in platemaking
INVENTOR (S):
                         Ide, Koji; Fumiya, Shinichi; Sasa, Nobumasa;
                         Watabe, Manabu
PATENT ASSIGNEE(S):
                         Mitsubishi Chemical Industries Co., Ltd.,
```

Japan; Konica Co.

SOURCE: Jpn. Kokai Tokkyo Koho, 10 pp.

CODEN: JKXXAF

DOCUMENT TYPE: LANGUAGE:

Patent Japanese

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

KIND PATENT NO. DATE APPLICATION NO. DATE JP 62291634 A2 19871218 JP 1986-134536 1986 0610 PRIORITY APPLN. INFO.: JP 1986-134536 1986

0610

AB A pos.-working color photoimaging sheet provides a color proof whose image quality is as good as that from printing. The color image-forming layer of the color photoimaging sheet is given the following components to improve its sensitivity. It contains (a) a compound such as 2-trichloromethyl-5-[β-(2-benzofuryl)vinyl]-1,3,4-oxadiazole that generates an acid upon irradiation with an actinic ray and (b) a compound such as an  $\alpha$ -thioglyceroltrimethyl orthoformate copolymer having ≥1 bond dissociable by the acid. The color photoimaging sheet is prepared by laminating on a transparent support a heat-fusing layer and a color image-forming layer. A color image-forming process includes imagewise exposure of the photoimaging sheet, development, and transferring the resultant color image to a receptor sheet by applying heat and pressure and peeling off the transparent support.

IT 115324-68-0

> (color proof for platemaking from photoimaging composition containing)

RN 115324-68-0 HCAPLUS

CN 1,2-Propanediol, 3-mercapto-, polymer with trimethoxymethane (9CI) (CA INDEX NAME)

CM 1

CRN 149-73-5 CMF C4 H10 O3

OMe MeO-CH-OMe

> CM 2

CRN 96-27-5 CMF C3 H8 O2 S

```
OH
HS-CH_2-CH-CH_2-OH
     ICM G03C001-00
IC
     ICS B41M005-26; G03C005-00; G03C005-24; G03F003-10
     74-6 (Radiation Chemistry, Photochemistry, and
CC
     Photographic and Other Reprographic Processes)
IT
     Printing plates
        (color proofing in production of, photoimaging compns.
        for)
IT
     Photoimaging compositions and processes
        (color, pos.-working, for color proofing in platemaking)
IT
     93641-24-8 115324-68-0 115324-69-1 115324-80-6
        (color proof for platemaking from photoimaging composition
        containing)
L21 ANSWER 21 OF 25 HCAPLUS COPYRIGHT 2006 ACS on STN
                         1967:444992 HCAPLUS
ACCESSION NUMBER:
                         67:44992
DOCUMENT NUMBER:
TITLE:
                         Xanthate decomposition in
                         acid media. IV. Cellulose xanthate;
                         dissociation equilibrium and kinetics of
                         decomposition
AUTHOR (S):
                         Tornell, Bertil
CORPORATE SOURCE:
                         Chalmers Tek. Hogskol., Goteborg, Swed.
SOURCE:
                         Svensk Papperstidning (1967), 70(9), 303-14
                         CODEN: SVPAAE; ISSN: 0283-6831
DOCUMENT TYPE:
                         Journal
LANGUAGE:
                         English
     cf. preceding abstract The decomposition of ripened and unripened
     samples of cellulose xanthate in acid medium was studied
     spectroscopically and the results showed that the intrinsic
     dissociation constant (K0) increased with decreasing number d. of unionized
     xanthate groups along the macromol. backbone. The limiting
     intrinsic pKa value obtained when the number d. of unchanged xanthate
     groups approached 0 was estimated as 1.0-1.2, indicating that
     cellulose xanthic acid is much stronger than Et xanthic acid.
     second order reaction rate constant (k) varied with the number d. of
     uncharged xanthate groups and the variation was of the same
     magnitude but of opposite sign to that of KO. This simultaneous
     variation was explained by assuming that the effective dielec.
     constant decreased with increasing number d. of unchanged xanthate
     groups. At low degrees of substitution, the product kKO was
     dependent on the composition of the reaction medium with
     variations similar to those observed with Et xanthate. Intramol.
     interactions exerted a stabilizing effect on the xanthate groups.
IT
     9032-37-5, Cellulose, hydrogen dithiocarbonate
        (dissociation equilibrium in acid and kinetics of
        decomposition of)
RN
     9032-37-5 HCAPLUS
CN
     Cellulose, hydrogen carbonodithioate (9CI) (CA INDEX NAME)
     CM
     CRN 9004-34-6
     CMF Unspecified
     CCI PMS, MAN
```

\*\*\* STRUCTURE DIAGRAM IS NOT AVAILABLE \*\*\*

CM 2

CRN 4741-30-4 CMF C H2 O S2

0 HS-C-SH

CC 43 (Cellulose, Lignin, Paper, and Other Wood Products) IT 9032-37-5, Cellulose, hydrogen dithiocarbonate (dissociation equilibrium in acid and kinetics of decomposition of)

L21 ANSWER 22 OF 25 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1964:433152 HCAPLUS

61:33152

DOCUMENT NUMBER:

ORIGINAL REFERENCE NO.: 61:5833g-h,5834a

TITLE:

Mechanism of capillary viscose-fiber formation. XX. Kinetics of regenerated

cellulose fiber formation. 3. The

decomposition of cellulose xanthate in acid Groebe, Anneliese; Jost, Hartmut; Klare,

Hermann

CORPORATE SOURCE:

Deut. Akad. Sci., Berlin

SOURCE:

AUTHOR (S):

Faserforschung und Textiltechnik (1964),

15(5), 193-201

CODEN: FSTXA7; ISSN: 0014-8628

DOCUMENT TYPE:

Journal

LANGUAGE:

Unavailable

cf. CA 61, 3248c. The effects of various parameters on the course of cellulose xanthate decomposition were followed by reaction-rate constant detns. The constant did not depend on the Hottenroth ripeness or on the initial degree of xanthation in the range of 29.0-48.7 studied. The constant increased with a decreasing fiber diameter; however, the effect of this parameter was almost negligible in fiber diams. above 1 mm. At those larger diams., the rate-determining factor was diffusion rather than cellulose xanthate decomposition The relations between reaction-rate constant (R) and temperature (T) were expressed by the equation RT2 = RT1 exp[(A)/B(T2 - T1), where B represents the universal gas constant (1.987 cal./degree); T, the absolute temperature (°K.); A, 17 ± 2 kcal./mole for pure cellulose xanthate solns. and model filaments and 6  $\pm$  0/5 kcal./mole for alkaline Na cellulose xanthate filaments. xanthate, both in the form of an emulsion and of a filament, decompose slower in the 0-5 pH range than the corresponding Na cellulose xanthate. The relations between decomposition rate and pH are expressed in the equation log R = CpH + d, where C and d are consts., depending on the composition of the xanthate. The filament-formation process consists of neutralization of the alkali in the model filament, acidification of the model filament, and regeneration of the cellulose xanthate at the pH of the spinning bath. The latter stage is the rate-determining step in filament formation. The above findings applied also to capillary viscose filaments.

```
IT
     9032-37-5, Cellulose xanthate
        (decomposition degradation or hydrolysis of, in acids in filament
        formation in spinning)
     9032-37-5 HCAPLUS
RN
CN
     Cellulose, hydrogen carbonodithioate (9CI) (CA INDEX NAME)
     CM
     CRN 9004-34-6
         Unspecified
     CMF
     CCI PMS, MAN
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
     CM
          2
     CRN 4741-30-4
     CMF C H2 O S2
   0
HS-C-SH
     47 (Textiles)
CC
IT
     Rayon
        (spinning of, filament formation by xanthate decomposition
        in acid in)
TΨ
     Cellulose xanthates, sodium salt
     Cellulose xanthates, zinc salt
        (formation and decomposition in acid in filament
        formation)
IT
     Zinc, with cellulose xanthates
        (formation and decomposition in acids in
        filament formation)
IT
     9032-37-5, Cellulose xanthate
        (decomposition degradation or hydrolysis of, in acids in filament
        formation in spinning)
L21 ANSWER 23 OF 25 HCAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                         1961:147201 HCAPLUS
DOCUMENT NUMBER:
                         55:147201
ORIGINAL REFERENCE NO.: 55:27897i,27898a-b
                         Formation mechanism of regenerated cellulose
TITLE:
                         fibers. XIII. Fiber formation reactions of
                         capillary viscose fibers
AUTHOR (S):
                         Grobe, Anneliese; Maron, Reinhard; Klare,
                         Hermann
CORPORATE SOURCE:
                         Deut. Akad. Wiss., Berlin
SOURCE:
                         Faserforschung und Textiltechnik (1961), 12,
                         196-207
                         CODEN: FSTXA7; ISSN: 0014-8628
DOCUMENT TYPE:
                         Journal
LANGUAGE:
                         Unavailable
     cf. CA 55, 21597g. Acid penetration in capillary viscose
     filaments emerging from the spinnerets (bromocresol purple being
     used as indicator) was studied. Upon completion of the
     acid penetration, the decomposition of the Na
     cellulose xanthate was complete and the only reaction that took
```

place thereafter was neutralization of any alkali present. neutralization point was a quant. measure of the H+ activity of the spinning bath, regardless of its composition In a H2SO4/Na2SO4 spinning bath, the fiber cross-section was circular if the concentration was ≤2N NaHSO4; above this concentration, the cross-section resembled a folded ribbon. The differences in fiber cross-section were attributed to the shrinking effect of the spinning bath employed. All reactions pertinent to fiber formation were completed within 50 mm. from the spinneret orifice at normal spinning rates. 9032-37-5, Cellulose xanthate IT (sodium derivs., decomposition in acid penetration in spinning viscose filaments) 9032-37-5 HCAPLUS RN Cellulose, hydrogen carbonodithioate (9CI) (CA INDEX NAME) CN CM 1 CRN 9004-34-6 CMF Unspecified CCI PMS, MAN \*\*\* STRUCTURE DIAGRAM IS NOT AVAILABLE \*\*\* CM 2 CRN 4741-30-4 CMF C H2 O S2 0 HS-C-SH CC 25 (Dyes and Textiles) IT 9032-37-5, Cellulose xanthate (sodium derivs., decomposition in acid penetration in spinning viscose filaments) L21 ANSWER 24 OF 25 HCAPLUS COPYRIGHT 2006 ACS on STN 1960:84760 HCAPLUS ACCESSION NUMBER: DOCUMENT NUMBER: 54:84760 ORIGINAL REFERENCE NO.: 54:16139e-i,16140a-b TITLE: Kinetic studies on the decomposition of xanthates AUTHOR (S): Philipp, Burkart; Fichte, Christa CORPORATE SOURCE: Deut. Akad. Wiss., Berlin SOURCE: Faserforschung und Textiltechnik (1960), 11, 118-24,172-9 CODEN: FSTXA7; ISSN: 0014-8628 DOCUMENT TYPE: Journal LANGUAGE: Unavailable The decomposing solns. consisted of aqueous 0.1 to 10M NaOH solns. for the strongly alkaline pH range and of buffered NaOH solns. for the pH 5-11 range. Initial xanthate concns. ranged from 0.01 to 0.1 mole/1. The decomposition kinetics were studied at 30 and 20°; and for ethylxanthate and O-Et monothiocarbonate also in the 65-85° temperature range. The xanthate (monothiocarbonate)

content in the reaction mixture and the sulfide (trithiocarbonate)

content in the end product were determined at such intervals as to yield 8-10 individual results in the course of 2-3 times of the resp. half-life time. The decomposition of xanthates in an acid medium was characterized by a rapid hydrolytic formation of the free xanthic acid which, in turn, decomposed under the formation of CS2, the only S-containing decomposition product. At a constant pH the reaction was of the 1st order. The same scheme applied to the decomposition of Et monothiocarbonate. In a strongly alkaline (above 5M NaOH) medium 2 xanthate ions and an OH ion formed an orthoxanthate in low concentration which, in turn, was split by a 2nd OH ion. On increasing the NaOH concentration from 7 to 10M the reaction rate constant increased considerably. Water did not appear to have participated in the decomposition mechanism. It was assumed that the orthoxanthates decompose into dithiocarbonates 1st; the latter then transform into sulfide through monothiocarbonate. Trithiocarbonate formed only at high alkaline concns. from cellulose xanthate. The absolute values of the reaction rate consts., in 7N NaOH at 30° varied from 0.022 to 1.0/hr. in the order: Et < 1,4-butanediol < cellulose < 1,3-butanediol < glycol < glycerol xanthate. At medium and low alkalinity the decomposition proceeds in various ways according to the composition of the xanthate. Et xanthate decompose through the slow hydration of the C:S bond; the primary product rapidly transforms into dithiocarbonate. The decomposition of polyalcs. was characterized by a secondary mechanism between the xanthate ions and water in which the adjoining free OH group also participated; the latter increased the reaction rate. This effect of the OH groups was even more pronounced in the case of glycerol and glycol xanthates, where, in the course of decomposition, any excess of free glycerol or glycol, resp., could cause rexanthation through the anionization of the vicinal OH groups. In cellulose xanthate and xanthates of simple carbohydrates the vicinal free OH groups facilitated the transformation with water, the transformation with OH ions playing a secondary function. A stability maximum of cellulose xanthate at 2-3N NaOH alkalinity was attributed to an equilibrium between a slowly proceeding reaction of xanthates and OH- (this reaction decelerates with a decreasing NaOH concentration) and a reaction of xanthate groups and water (the rate increasing with decreasing NaOH concentration). The greater stability, under identical conditions, of C6 xanthates, as compared with C2-3 xanthates, was attributed to the action of vicinal undissocd. OH groups in the former. 9032-37-5, Cellulose xanthate

IT

(decomposition of)

RN 9032-37-5 HCAPLUS

Cellulose, hydrogen carbonodithioate (9CI) (CA INDEX NAME) CN

CM

CRN 9004-34-6 CMF Unspecified CCI PMS, MAN

\*\*\* STRUCTURE DIAGRAM IS NOT AVAILABLE \*\*\*

CM

CRN 4741-30-4 CMF C H2 O S2

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HS-C-SH
CC
     2 (General and Physical Chemistry)
IT
     Hydroxyl group
        (effect of vicinal, on decomposition of xanthic
        acid derivs. of polyhydric alcs.)
IT
     140-89-6, Xanthic acid, ethyl-, potassium salt 9032-37-5
     , Cellulose xanthate
                           22410-33-9, Xanthic acid, 2-hydroxyethyl-,
                      35832-93-0, Carbonic acid, thio-, O-ethyl ester,
     potassium salt
                      57516-89-9, Xanthic acid, (2,3-dihydroxypropyl)-,
     potassium salt
     potassium salt
                      116600-70-5, Xanthic acid, 4-hydroxybutyl-,
     potassium salt
                      116600-72-7, Xanthic acid, 3-hydroxybutyl-,
                      116603-95-3, Xanthic acid, D-glucosyl-, potassium
     potassium salt
            117042-93-0, Xanthic acid, (3-hydroxy-1-methylpropyl)-,
     salt
                      860178-56-9, Xanthic acid, [2-hydroxy-1-
     potassium salt
     (hydroxymethyl)ethyl]-, potassium salt
        (decomposition of)
L21 ANSWER 25 OF 25 HCAPLUS COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER:
                         1941:21519 HCAPLUS
DOCUMENT NUMBER:
                         35:21519
ORIGINAL REFERENCE NO.: 35:3435c-q
TITLE:
                         The effect of salts of heavy metals on the
                         velocity of decomposition of cellulose
                         xanthate
AUTHOR (S):
                         Pakshver, A.; Frolov, S.; Karegina, N.
SOURCE:
                         Trans. Inst. Chem. Tech. Ivanovo (U. S. S. R.)
                         (1940), (No. 3), 178-80
DOCUMENT TYPE:
                         Journal
LANGUAGE:
                         Unavailable
AB
     The object of the expts. was to amplify the results obtained by
     Danilov and Gintse (C. A. 33, 5179.7) and to verify the decomposition
     velocity of Na, Ca, Zn, Ni and Cu cellulose xanthates. A 1-mm.
     layer of ordinary viscose was placed on a glass plate, coagulated
     in concentrated NaCl solution for 20 min., washed on the glass with
concentrated
    NaCl solution to complete removal of impurities (sulfides,
     thiocarbonates, etc.) and treated for 10 min. with a concentrated solution
     of the corresponding salt (CaCl2, NaCl, CuCl2, ZnCl2).
     Preliminary expts. showed that a 10-min. reaction is sufficient.
     The film was decomposed with N H2SO4 for a definite time (with a
     stop watch) and immediately immersed in a 10% NaOAc solution, after
    which it was immersed for 5 min. in 20 ml. of AcOH and 10 ml. of
     0.1 N I solution The amount of AcOH and the time of the reaction were
    determined by a blank experiment  The excess I was titrated with thiosulfate
     and the percentage of xanthate decomposed calculated The velocity of
     the decomposition of xanthate does not increase with aging of the
    viscose, but decreases slightly. A decrease takes place also with
     the increase of the ratio \alpha-cellulose:NaOH in viscose. The
    decomposition velocity is determined by the colloidal properties of the film
    of the cellulose. The older xanthates form on decomposition
    with acid a denser surface film of the regenerated
    cellulose, which retards the decomposition This explains also the
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effect of the composition of viscose on the decomposition velocity of the xanthate. NiSO4 retards the decomposition of the xanthate to a

smaller extent than does ZnSO4. Therefore ZnSO4 cannot replace

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NiSO4 in the precipitating bath.
     9032-37-5, Cellulose xanthate
IT
        (decomposition of, effect of salts of heavy metals and compn
        . of viscose on)
     9032-37-5 HCAPLUS
RN
    Cellulose, hydrogen carbonodithioate (9CI) (CA INDEX NAME)
CN
    CM
          1
         9004-34-6
    CRN
     CMF
         Unspecified
    CCI PMS, MAN
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
    CM
          2
    CRN 4741-30-4
    CMF C H2 O S2
   0
HS-C-SH
    23 (Cellulose and Paper)
CC
IT
    Viscose
        (effect of composition of, on decomposition velocity of
        cellulose xanthate)
IT
    Reaction kinetics and (or) velocity
        (of cellulose xanthate decomposition, effect of heavy-metal salts
        and composition of viscose on)
IT
     9032-37-5, Cellulose xanthate
        (decomposition of, effect of salts of heavy metals and compn
        . of viscose on)
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